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(71) Applicant: CYTOKINETICS, INC. [US/US]; Suite 2, 280 East Grand Avenue, South San Francisco, CA 94080 (US).			
(72) Inventors: SABRY, James, H.; 52 Buena Vista Terrace, San Francisco, CA 94117 (US). ADAMS, Cynthia, L.; 615 Georgia Avenue, Palo Alto, CA 94306 (US). VAISBERG, Eugeni, A.; 647 Pegasus Lane, Foster City, CA 94404 (US). CROMPTON, Anne, M.; 2 Bellair Place, San Francisco, CA 94133 (US). BLUM, Robert, I.; 17 Shoreview Avenue, San Francisco, CA 94121 (US). OESTREICHER, Donald, R.; 904 Old Town Court, Cupertino, CA 95014-4024 (US). SIGAL, Nolan, H.; 941 Berry Avenue, Los Altos, CA 94024 (US).			

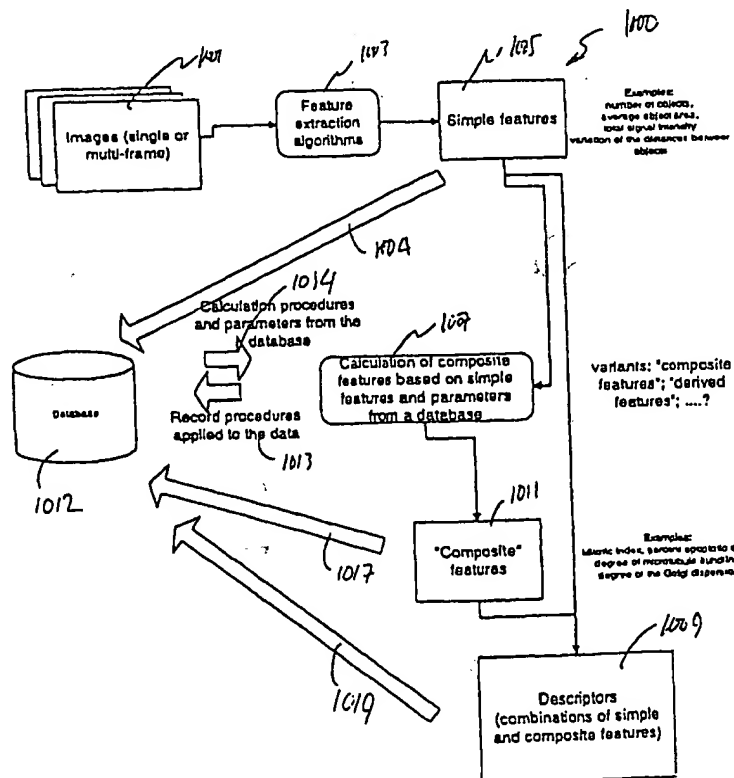
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(57) Abstract

Techniques for using information technology in therapeutics or drug discovery. In an exemplary embodiment, techniques for determining information about the properties of substances based upon information about structure of living or non-living cells exposed to substances are provided. A method according to the present invention enables researchers and/or scientists to identify promising candidates in the search for new and better medicines or treatments using, for example, a cellular informatics database. The present invention further teaches a system for acquiring knowledge from cellular information. The system has a database 1012 comprising a database management module ("DBMS"). The system also has a variety of modules, including a population module coupled to the DBMS for categorizing and storing a plurality of features (e.g., cell size, distance between cells, cell population, cell type) from an image acquisition device into the database. The system has a translation module coupled to the DBMS for defining a descriptor from a set of selected features from the plurality of features. In a specific embodiment, the descriptor is for a known or unknown compound, e.g., drug. A prediction module is coupled to the DBMS for selecting one of a plurality of descriptors from known and unknown compounds from the database based upon a selected descriptor from a selected compound. The selected compound may be one that is useful for treatment of human beings or the like.



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PATENT APPLICATION
METHOD AND APPARATUS FOR
PREDICTIVE CELLULAR BIOINFORMATICS
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10 computer codes, which may be used to implement aspects of the present invention. Assignee of the present invention reserves all rights with respect to these codes and provides notice herein. Notice is hereby given © Cytokinetics, Inc. 1999.

BACKGROUND OF THE INVENTION

15 The present invention provides techniques for information management using a database platform. More particularly, the present invention provides a system including computer code that couples to a database device. The system provides for image capturing of living, dead, or fixed cells or cell fractions used to identify information about substances used on the cells or information about the cells themselves. Accordingly, the present invention can enable researchers and
20 scientists to identify promising candidates in the search for new and better medicines, for example, in drug discovery and development. The principles enumerated herein may, with equal facility, be applied to other applications, including but not limited to use in environmental applications such as determining chemical toxicities and other non-pharmaceutical toxicology uses.

25 For a long time, researchers in the pharmaceutical field have sought for better ways of searching for substances possessing properties that make them suitable as medicines. In the early days, researchers generally relied upon extracts from plants, dyes, and microbiological extracts for such substances. Examples of such substances include the pain reliever aspirin, the anti-cancer drug paclitaxel (brand
30 name TaxolTM), and the heart medication called digoxin. The number of useful medicines has generally been limited.

Purified substances having desirable bio-active properties are also often difficult to discover. Advances in traditional organic chemistry and more recently the rapid chemical synthesis methods often referred to as combinatorial chemistry have increased the number of compounds that researchers test for biological activity. Originally, substances were often initially tested on animals or humans to determine their biological activity. While results from such tests may identify a good drug candidate, they are often time consuming and costly, thus a limited number of substances can be tested. Therefore, pharmaceutical companies have turned to testing their ever-increasing libraries of substances against isolated proteins (drug targets) in biochemical assays that can be carried out at high throughput and low cost. It should be noted that the substances need to be tested in numerous protein tests, each customized for a particular drug target. Therefore, although each protein test may be run at a high-throughput, the design of multiple protein tests can be time-consuming. Substances deemed promising based on results from the protein tests are then tested in lower throughput cellular and animal tests.

There have been some attempts to use image acquisition techniques to screen a large number of substances based upon biological cell information. One such attempt is described in International Application No. WO 98/38490 in the names of Dunlay, et al. Dunlay et al. generally describes a conventional image acquisition system. This conventional system collects and saves images based on certain criteria that are predefined, not on a fixed area of an imaging surface. Additionally, the conventional system has poor lighting design, which makes image processing for multiple cells difficult. Furthermore, the conventional system is not designed for capturing, populating and utilizing a large database design. The conventional system is designed for customized cellular assays, not as a tool for generation of a cellular informatics database. Without such database capabilities the conventional system cannot be used for screening, analyzing, and comparing large quantities of cells from multiple experiments on multiple days in a predictive, efficient and cost effective manner.

What is needed is a rapid assay to assess the activity of compounds against multiple drug targets simultaneously in a cellular context. What is also needed are techniques for finding the effects of substances on cell function based upon searching and analyzing cellular information.

SUMMARY OF THE INVENTION

According to at least one embodiment of the present invention, techniques for determining information about effects of potential substances on cells are provided. In another exemplary embodiment, the present invention provides a novel system including hardware, computer codes, user interfaces, and a database for acquiring, storing and retrieving cellular and substance information. The cells can include living, dead, or fixed cells or fractions of cells. The present invention enables, *inter alia*, researchers and/or scientists to identify promising candidates in the search for new and better medicines or treatments using, for example, a cellular informatics database.

According to the present invention, a computer program for identification and verification of biological properties of substances can include code that causes a sample of a substance to be administered to a cell. The code determines one or more features for two or more cell components, or markers, in the presence of the substance. The code can form one or more descriptors from the features. Descriptors can be formed by combining features of two or more cell components as identified using the markers. The code can then search one or more descriptors obtained from prior administered substances upon cells in order to locate descriptors having a relationship to the descriptors noted for the substance under study. The code predicts properties of the administered substance based upon the properties of the prior administered substances using the relationship between the descriptors. The code can provide for identifying properties of substances based upon effects on cell characteristics. Candidate drug mechanisms of action, potency, specificity, pharmacodynamic, and pharmacokinetic parameters, toxicity, and the like can be used as substance properties.

In a specific embodiment, the present invention provides a system for acquiring knowledge from cellular information. The system has a database comprising a database management module ("DBMS"). The system also has a variety of other modules, including a population module that is coupled to the DBMS and serves to categorize and store a plurality of features (including but not limited to cell size, distance between cells, cell population, as well as sub-cellular features such as organelle location, protein location and sub-cellular constituent location and

movement) from an image acquisition device into the database. The system has a translation module coupled to the DBMS for defining a descriptor from a set of selected features from the plurality of features. In a specific embodiment, the descriptor is for a known or unknown compound, e.g., drug. A prediction module is coupled to the DBMS for selecting one of a plurality of a descriptors from known and unknown compounds from the database based upon a selected descriptor from a selected compound. The selected compound may be one that is useful for treatment of human beings or the like.

In a specific embodiment, the present invention provides a system for populating a database with cellular information. The system includes a cell holder (e.g., multi-well plate, chip, microfluidic assembly, or other cell chamber) comprising a plurality of sites in a spatial orientation. Each of the sites is capable of holding a plurality of cells to be imaged. Note – the light guide is one embodiment, but we don't want to be limited to it.

According to one embodiment, the present system also has an illumination apparatus including a liquid light guide operably coupled to the imaging device for highlighting the plurality of cells in a relatively even spatial manner for image capturing and measurement purposes. Still further, the liquid light guide allows sub-elements (e.g., filter, lamp) of the illumination apparatus to be placed at a remote location to prevent mechanical interference of the cell holder during image capturing. Alternative lighting methodologies may, with equal facility, be implemented.

The system also has an image-capturing device (e.g., charge coupled device camera, translation stage, shutter, microscope, software, shutter control) coupled to a computing device (e.g., computer, network computer, work station, analog computing device, on-board image-processor, and laptop). The image-capturing device is adapted to capture at least one image in at least one of the plurality of sites. One some embodiments, multiple images can be captured, where each image represents a different cell component (or portion). The image-capturing device can be adapted to convert the image into a digital representation, which highlights the feature or features of the one site.

A database storage device (e.g., relational database, object oriented database, mixed object oriented database) includes a database management element. The

database is coupled to the image capturing device. In a specific embodiment, the present system includes modules for feature extraction, generation of descriptions, and data preparation and analysis.

In a specific embodiment, the present invention provides a novel
5 system for determining an effect of a manipulation of a cell using one or more image frames. The system has a plate comprising a plurality of sites in a spatial orientation. Each of the sites is capable of holding a plurality of cells to be imaged. The system also has an image capturing device to capture a plurality of images of at least one site from the plurality of sites. The image capturing device is coupled to the computing
10 device. The system also has an image processing device to combine the plurality of images of at least one site or plurality of sites. The image processing device is operably coupled to the plate. An image processing device is also included. The image processing device can be adapted to form a digitized representation of the plurality of images from the site or plurality of sites. Furthermore, the system has a
15 database storage device comprising a database management element. The database can be adapted to retrieve the descriptor or descriptors of the plurality of features from the computing processing device and storing them in a selected manner.

In a specific embodiment, the present invention provides a system for capturing cellular information. The system also has an image acquisition system
20 comprising a charged coupled device camera adapted to capture an image of a plurality of manipulated cells in various stages of the cell cycle. The stages of the cell cycle are currently understood to include interphase, G0 phase, G1 phase, S phase, G2 phase, M phase, prophase, prometaphase, metaphase, anaphase, and telophase. The principles of the present invention specifically contemplate the application thereof on
25 additional cell cycle stages when and if they are identified.

An optical source is coupled to the image acquisition system for highlighting the plurality of manipulated cells in the various stages of the cell cycle. The illumination apparatus provides for an acquisition of the image of the plurality of manipulated cells. In a specific embodiment, the illumination apparatus has a liquid
30 light guide coupled to a light source at a remote location.

A variety of user interfaces are utile for accessing the several features of the present invention. Those having ordinary skill in the art will appreciate that different user interfaces may be required to support different research scenarios. The

present invention specifically contemplates the utilization of a wide variety of user interfaces.

Numerous benefits are achieved by way of the present invention over conventional techniques. The present invention can provide techniques for predictive cellular bioinformatics that can streamline a number of important decisions made in the drug discovery industry. The present invention can be implemented using off the shelf hardware including databases. In other aspects, the present invention can find useful information about substances as well as cells or portions of cells. Furthermore, the present invention can acquire more than one feature using more than one manipulation. Moreover, the present invention can provide information about a wide variety of cellular information that is not conventionally available. This information includes information about different cell components, e.g., nuclei and Golgi apparatus. Still further, the present invention provides an automated or semi-automated technique for acquiring images and populating a database. The present database can be combined with others such as genomics, and the like. Moreover, the present invention can be implemented to predict, *inter alia*, a mechanism of action, toxicity, target validation, and pre-clinical disease model.

A further understanding of the nature and advantages of the invention herein may be realized by reference to the remaining sections of the specification and the attached drawings.

BRIEF DESCRIPTION OF THE DRAWING

For more complete understanding of the present invention, reference is
5 made to the accompanying Drawing in the following Detailed Description of the
Invention. In the drawing:

Fig. 1 is a simplified system diagram according to an embodiment
according to the present invention;

Figs. 1A-1B are more detailed diagrams of database systems according
10 to embodiments of the present invention;

Fig. 2 is a simplified block diagram according to an alternative
embodiment according to the present invention;

Figs. 3-6 are simplified diagrams of system elements according to
embodiments of the present invention;

15 Figs. 7A-7K illustrate representative block diagrams of simplified
process steps in a particular embodiment according to the present invention;

Fig. 8A-8F illustrate representative quantified descriptors of effects of
manipulations on images of cells in a particular experiment;

Fig. 9 illustrates example images for different types of morphologies in
20 a particular experiment;

Fig. 10 illustrates a distribution of various morphologies in a cell
population responsive to drug concentration in a particular experiment;

Fig. 11 illustrates a graph of quantified features of effects of
manipulations on cells in a particular experiment;

25 Fig. 12 illustrates effects of external agents on cells in a particular
experiment;

Fig. 13 illustrates 4 panels for each marker for a plurality of A549 cells
in a particular experiment;

Fig. 14 illustrates 4 panels for each marker for a plurality of OVCAR-3
30 cells in a particular experiment;

Fig. 15 illustrates 4 panels for each marker for a plurality of OVCAR-3
cells at 20x in a particular experiment;

Fig. 16 illustrates 4 panels for each marker for a plurality of OVCAR-3 cells at 40x in a particular experiment;

Fig. 17 illustrates a representative input for a morphometric analysis program in a particular embodiment according to the present invention; and

5 Figs. 18-19 illustrate examples of the generation of pseudo-sequences and clustering in a particular embodiment according to the present invention.

Fig. 20 is a block diagram for a first research scenario;

Fig. 21 is a block diagram for a second research scenario; and

Fig. 22 is a block diagram for a third research scenario.

10 Reference numbers refer to the same or equivalent parts of the invention throughout the several figures of the Drawing.

DETAILED DESCRIPTION OF THE INVENTION

According to the present invention, techniques for determining information about manipulated cells or substances based upon living, fixed, or dead cell structures or portions of cells are provided. In an exemplary embodiment, the present invention provides a novel system including computer codes coupled to a database and user interfaces for acquiring, storing and retrieving such information. Other embodiments provide a novel image capturing system for providing digitized representations of live and dead cell structures or the like.

Fig. 1 is a simplified system diagram 10 of a cellular knowledge-based system according to an embodiment of the present invention. This diagram is merely an example and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. The present system 10 includes a variety of elements such as a computing device 13, which is coupled to an image processor 15 and is coupled to a database 21. The image processor receives information from an image capturing device 17, which image processor and image capturing device are collectively referred to as the imaging system herein. The image capturing device obtains information from a plate 19, which includes a plurality of sites for cells. These cells can be biological cells that are living, fixed, dead, cell fractions, cells in a tissue, and the like. The computing device retrieves the information, which has been digitized, from the image processing device and stores such information into the database. A user interface device 11, which can be a personal computer, a work station, a network computer, a personal digital assistant, or the like, is coupled to the computing device.

Fig. 1A is a simplified diagram of a database system 1000 according to an embodiment of the present invention. This diagram is merely an example and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize many other variations, modifications, and alternatives. Database system 1000 includes a variety of techniques for processing images from biological cells, e.g., fixed, living, and dead cells, and cell portions. As shown, images are acquired 1001. These images can be from a single frame or multiple frames. As merely an example, an image processing system may analyze such images. One example of

such an image processing system is described below, but should not be construed as limiting certain claims.

In a specific embodiment, cell samples are manipulated using a compound (e.g., substance, drug). The cell samples are imaged for a simple portion or portions, e.g., manipulated cell substructure, manipulated spatial feature of cell, cell density. Image processing techniques are used to extract the feature or features from the image or images. The features can be an independent or a dependent set of cell characteristics (which may be predominately visual) including, for example, count, area, perimeter, length, breadth, fiber length, fiber breadth, shape factor, elliptical form factor, inner radius, outer radius, mean radius, equivalent radius, equivalent sphere volume, equivalent prolate volume, equivalent oblate volume, equivalent sphere surface, average intensity, total intensity, optical density, radial dispersion, texture difference, and others. Each of these features corresponds to a similar manipulation by a compound. Each manipulation forms a new set of features, which are identifiable to the compound. Once each set of features has been extracted, the feature set is populated into a database. Accordingly, the database includes many sets of features, where each set corresponds to a different manipulation for a selected cell. Each set of features corresponding to a manipulation provides a descriptor, which is also stored in the database. The descriptor is a "finger print" including each feature for the manipulation. Each descriptor may be unique, or may have similarities to other descriptors or may even be the same as other descriptors for known and unknown manipulations.

The present system retrieves features, which we define as simple features herein, and forms composite features from them. More than one feature can be combined in a variety of different ways to form these composite features. In particular, the composite feature can be any function or combination of a simple feature and other composite features. The function can be algebraic, logical, sinusoidal, logarithmic, linear, hyperbolic, statistical, and the like. Alternatively, more than one simple feature can be combined in a functional manner (e.g., arithmetic, algebraic). As merely an example, the composite feature equals a sum of feature 1 and feature 2, where these features correspond to the same manipulation. Alternatively, the composite feature equals feature 1 divided by feature 2. Alternatively, the composite feature equals feature 1 minus feature 2. Alternatively,

the composite feature equals a constant times feature 1 plus feature 2. Of course, there are many ways that the composite feature can be defined. The present system also stores 1017 these features in the database. The composite features can also be further combined with simple features. Once these features are defined as descriptors, they are stored 1019 in the database.

Fig. 1B is a simplified diagram of a database system engine 2000 according to an embodiment of the present invention. This diagram is merely an example and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize many other variations, modifications, and alternatives. The engine can be implemented into the present database for populating, searching, and predicting compound or cell characteristics. As merely an example, engine 2001 includes an input/output module 2008. The input/output module is used to input and output information from the database. The information includes, among others, a plurality of feature sets, which correspond to many manipulations. Additionally, the information includes descriptors, which each corresponds to a set of features from the manipulation. The database also has a population module, which is used to configure the features based upon an entity relationship, which has been predetermined.

The database engine also has other modules. In particular, the database has a transcription module, which transfers a preselected set of features and creates a descriptor from them. The transcription module can be used to take a known compound, which has features, to transcribe them into a descriptor. Alternatively, the transcription module can be used to take an unknown compound, which has features, to transcribe them into a descriptor. These descriptors are provided into the database for subsequent use. Finally, the database engine has a prediction module, which can be used to potentially predict a property (e.g., mechanism of action) of an unknown compound. Here, the unknown compound is provided with a descriptor, but the property of the compound is unknown. In one embodiment, the prediction module compares a descriptor of an unknown compound with the many descriptors of known compounds, which were in the populated database. Depending upon the matching criteria, the prediction module will attempt to uncover one or more descriptors of known compounds. Once the prediction module finds the descriptors of the known compounds based upon the descriptor for the unknown compound, it identifies a potential property of such unknown compound for analysis and review. Here, it is

believed that certain features of the known compound, which are similar to those features of the unknown compound may uncover a property to the unknown compound. Details of the present software engine are described more fully below.

Fig. 2 is a simplified block diagram 20 of a cellular knowledge-based system according to an alternative embodiment of the present invention. This diagram is merely an example and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. Like reference numerals are used in the present diagram as the previous diagram for easy cross-referencing, but are not intended to be limiting in any manner. The present diagram 20 includes a variety of elements such as a processor 13 or computing device coupled to a database 11. The processor can be used for retrieving and storing information from the database. The system also includes a plurality of system elements, such as a cleaner 23, a dispenser 25, and an image capturing system 27, which are also coupled to the database in some embodiments. These elements can be coupled to each other through a network or the like. As merely an example, the network can be a NetWareTM network from Novell Corporation or an internet network or the Internet but can also be others and any combination thereof. The system also has an output device 31, which can be used to output information from the database, processor, or other system elements. Details of these elements are described more fully below in reference to the Figs.

Figs. 3-5 are simplified drawings of system elements according to embodiments of the present invention. These diagrams are merely examples and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. As merely an example, Fig. 3 is a simplified diagram of a processor or computing device 13. The computing device 13 includes a bus 112 which interconnects major subsystems such as a central processor 114, a system memory 116 (e.g., random access memory), an input/output ("I/O") controller 118, an external device such as a display screen 124 via a display adapter 126, a keyboard 132 and a mouse 146 via an I/O controller 118, a SCSI host adapter (not shown), and a floppy disk drive 136 operative to receive a floppy disk 138.

The computing device has other features. Storage Interface 134 may act as a storage interface to a fixed disk drive 144 or a CD-ROM player 140 operative

to receive a CD-ROM 142. Fixed disk 144 may be a part of computing device or may be separate and accessed through other interface systems. A network interface 148 may provide a direct connection to a remote server via a telephone link or to the Internet. Network interface 148 may also connect to a local area network ("LAN") or other network interconnecting many computer systems. Many other devices or subsystems (not shown) may be connected in a similar manner. Also, it is not necessary for all of the devices shown in Fig. 3 to be present to practice the present invention, as discussed below. The devices and subsystems may be interconnected in different ways from that shown in Fig. 3. The operation of a computer system such as that shown in Fig. 3 is readily known in the art and is not discussed in detail in this application. Computer code to implement the present invention, may be operably disposed or stored in computer-readable storage media such as system memory 116, fixed disk 144, CD-ROM 140, or floppy disk 138. The computer code can be organized in terms of processes or modules, depending upon the application. That is, the computer code can include a prediction module, a translation module, or other modules to carryout the functionality described herein, as well as others.

Figs. 4 and 5 are simplified diagrams of an imaging system 200 according to an embodiment of the present invention. As shown, the imaging system 200 includes a variety of features such as housing 203, which holds a stage assembly 204. The stage assembly includes an x-stage movement element 206, which is along an x-direction, and a y-stage movement element 207, which is along a y-direction. The imaging system also includes a z-direction movement element, which is perpendicular to the x-y plane. The z-direction movement motor can be attached to the stage, or to the objective nosepiece by way of the microscope housing, or as an external motor between the objective and the microscope housing. The stage can align in any one of the directions to an accuracy of one micron and less, or one-half micron and less, or one-quarter micron and less, depending upon the embodiment.

The stage holds a plate 202 or cell holder, which houses one of a plurality of samples. The plate includes a spatial array 209 of process sites. Each of the process sites can include a plurality of cells and solutions depending upon the embodiment. Each of the sites can carry a sufficient amount of solution to prevent substantial evaporation of the sample during processing in some embodiments. In embodiments for large scale analysis, the plate includes at least 96 sites, or more than

or equal to 384 sites, or more than or equal to 1,536 sites. The plate bottom is transparent and thin, which allows light to pass through the sample. Additionally, the plate is made of a suitable chemical resistant material. As merely an example, the plate can be either a 96, or 384, or 1536 or other formats from places such as Becton Dickinson of Franklin Lakes, NJ, or Corning Science Products of Corning, NY. In a preferred embodiment, the plate is a Corning Costar black-walled 96 well plate catalog #3904 from Corning Science Products of Corning, NY, but should not be limited to these in some applications, but can be others.

Also shown is the condenser for the microscope 201, which can be used to collect phase, DIC, or bright field images of the cells. Images resulting from the illumination of the samples to fluorescence, phase, DIC, or bright field techniques are collected using an image capturing device 208, which captures an image or images of cells from the plate. In a specific embodiment, the microscope is an inverted configuration with the objectives on the bottom of the plate and the condenser disposed overlying an upper surface of the sites, while the image capturing device underlies the sites. Images captured by the imaging device, whether analogue or digital, are viewed by a monitor or other devices. The image capturing device can be any camera assembly such as a charge coupled device camera, which is known as a CCD camera, or other high resolution camera capable of capturing images from the sites. In a specific embodiment, the camera is an interline CCD camera which does not require an external shutter.

In a specific embodiment, the present imaging system can be any suitable unit that is flexible for automated image collection using multi-well plastic plates. The imaging system also should be adapted to collect high-resolution images of cells on plastic or glass plates, cell growth chambers, or coverslips. The system also can be used for imaging multiple cell markers in multiple imaging conditions. To accomplish this, the microscope system has a variety of elements such as a light source, a motorized excitation filter wheel and shutter, x-y-z-motorized stage, excitation and emission filters, Fluor phase and DIC objectives, motorized objective nosepiece, dichroic filters, motorized dichroic filter cubes, phase and DIC rings and prisms, CCD camera, and software control. As merely an example, the present imaging system can have components such as those listed in the Table below.

DESCRIPTION	MAKER	MODEL
Microscope	Zeiss	100M
(x-y) motorized stage	Prior	
Xenon lamp	Sutter	Lambda
Filter wheel	Sutter	Lambda-10
Microtitre Plate holder	Prior	500-H223R
Isolation Table	Kinetic Systems	9101-24-85
Objective Spacers	Polytec PI	P-721.90
Camera	Hamamatsu	C47-95
Computer	IBM	IntelliStation
Software	Metamorph	v.4
Objectives	Zeiss	Achroplan 10x/0.25 LD-Achroplan 20x/0.4 LD-Achroplan 40x/0.6

Table: Image Acquisition System Elements

5 In a specific embodiment, the present system has the following capabilities, which are not intended to be limiting.

Image acquisition

1) Ability to automatically acquire multi-wavelength images from multiple sites on one multi-well plate, to sequentially name image files, and to log any
10 imaging parameter information with image files.

2) Ability to link images with a larger database/spreadsheet of information.

3) Ability to automatically collect multiple plates by interfacing the imaging system with a robotic arm.

15

X-Y control

1) Ability to place 96, 384, or 1536 well plates onto microscope stage and move to each well sequentially.

2) Ability to return to each well and collect another round of images (multi-site time-lapse) or ability to collect rapid time-lapse information at each well (time-lapse of many wells).

3) Ability to collect a low magnification image, automatically
5 determine features which may be of interest, automatically change the objective to a higher magnification, and collect high magnification images of a fixed number of those identified cells in the sample.

4) Ability to collect multiple frames in each site.

10 Z control

1. Ability to auto-focus with substantially minimal damage to biological specimen or fluorophore.

2. Ability to auto-focus rapidly.

15 The present embodiment of the imaging system is shown by way of Figs. 5A and 5B. These diagrams are merely examples and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. The present imaging system 40 includes a variety of elements such as a microscope 41, which is preferably an epi-fluorescent microscope,
20 but can be confocal, multiphoton, or hybrid types. The microscope includes elements 41A, the motorized Z-axis; 41B, the motorized dichroic filter cube holder; and 41C, the motorized objective nosepiece. In one embodiment, the microscope is a Model 100M made by Zeiss. The microscope communicates to computer 51 through control lines 73, 75, and 76. The imaging system also has camera 50 coupled to controller
25 50A and computing device 51, which oversees and controls operations of the elements of the imaging system.

The present microscope includes drivers for spatially moving a stage in two dimensions, including an x-direction, a y-direction, and moving the objective nosepiece in a z-direction in a Cartesian coordinate system. The z-direction
30 movement is provided using a fast z-motor, which can make z-direction adjustments within a predetermined time. The z-direction movement generally provides for focussing of the sample to the camera. The focussing occurs within the predetermined time of preferably ten seconds and less, or five seconds and less, or one

second and less, depending upon the embodiment. As merely an example, the z-motor or positioner can be a model PIFOC objective nanopositioner made by a company called Physik Instrumente of Waldbronn, Germany, but also can be others. The z-motor couples to computer 51 through line 63, which may also include a
5 controller. Depending upon the embodiment, a second z-motor 41A connected to the computer 51 by line 73 may be used to keep the z-motor 42 in the center of its travel. Alternatively, in other embodiments the stage could be provided with a z-motor allowing for movement of the stage in the z-direction.

The present stage also includes an x-y stage 43. The x-y stage moves
10 plate 59, e.g., 96 site, 384 site, 1536 site. The x-y stage moves plate in an x-y spatial manner. The stage has an accuracy or repeatability of about 1 micron and less, or about 2 microns and less. The stage can move in a continuous manner or a stepped manner. The stage also can move up to 30 mm/sec. or faster. The stage also can move 1 mm/sec. and less, depending upon the embodiment. The stage can also step
15 0.1 micron and less or 1 micron and less, as well as other spatial dimensions. The stage can be one such as a Proscan Series made by Prior Scientific of Rockland, MA but can also be others. The stage is controlled via control line 61 through controller 43A, which couples to computer 51 through control line 65.

The stage includes plate holder 44. The plate holder can hold a single
20 plate. In other embodiments, plate holder can also hold multiple plates. The plate holder can use mechanical, electrical, fluid, vacuum and other means for holding the plate or plates. The plate holder also is sufficiently stable for securing the plate. As merely an example, the plate holder is a Model 500-H223R made by Prior Scientific of Rockland, MA. In some embodiments, the plate holder may need adjustment in
25 the z-direction to provide for a desirable focus of a sample on a plate. In these embodiments, the plate holder is supported by spacers 45 or a plurality of stage pins, which mechanically elevate the plate holder in the z-direction. These pins are generally made of a suitable material for supporting such plate holder and also are sufficiently resistant to chemicals and the like.

30 In some embodiments, the entire imaging system is placed on an isolation table 49. The isolation table is disposed between the microscope and support structure. The isolation table is designed to prevent excessive vibration of the plate. The isolation table is made of a suitable material such as steel and honeycomb but can

be others. The table has a thickness of about 8 inches or preferably less than about 24 inches. In one embodiment, the table is Model 9101-24-85 made by Kinetic Systems of Boston, MA.

The imaging system also has a lamp or illumination assembly 62. The lamp assembly provides for a light source (See reference letter B) to a plurality of elements in the imaging system. For easy reading, the light path is defined by the dotted lines, which are not intended to be limiting. The lamp assembly has a variety of elements such as a Xenon lamp 46. The Xenon lamp provides light at about 320 to 700 nanometers (Prefocused). The Xenon lamp is 175 or 300 Watts. As merely an example, the lamp can be a Lambda Model made by Sutter Instrument Company of Novato, CA.

Referring to Fig. 5B, the lamp assembly also has a cold mirror 58, an excitation filter wheel 48, excitation filter(s) 55, and an excitation light shutter 57. As shown, light is derived from the Xenon lamp, reflects off of the cold mirror 58, traverses through the excitation filter or filters 55, and is controlled by the excitation light shutter 57. The lamp assembly has filter wheel 48, which houses one of a plurality of filters, including excitation filters. The shutter and filter wheel are controlled via control lines 67, which are coupled to a computer 51 or other type of computing device. The control lines 67 are coupled through controller 57A (for element 57) and controller 48A (for element 48) via control line 69 to computer 51.

Preferably, light traverses from the lamp assembly through a light guide 47 to illuminate features within the plate. The light guide is suitably selected to have a flexible member, which can be used to place lamp source at a remote location away from the imaging device. The flexible member substantially keeps any vibration from the lamp assembly away from the imaging device. In some embodiments, the member is at least 1 foot away from the imaging device. The light guide is a guide, which is a flexible hose-type sleeve. The sleeve is filled with a liquid such as an aqueous solution containing chloride or phosphate. A thin layer may be formed on the inside of the sleeve. The layer can be a containing tetrafluoroethylene and hexafluoropropylene, or containing tetrafluoroethylene and perfluoromethyl vinyl ether, or tetrafluoroethylene and perfluoropropyl vinyl ether. An example of such a light guide is described in International Application No. WO/98/38537 filed February 29, 1997, and assigned to NATH, Gunther. The liquid

light guide has less than about 30% transmission loss of the light at a remote location such as the imaging system.

Light is derived from the lamp assembly and directs off of filter 56, which directs the light upward. Filter 56 can be a dichroic and emission filter, as well as others. The light traverses through microscope nosepiece 41C, and traverses
5 through objective spacers 54. An objective 53 magnifies the light toward a predetermined point on the plate 59. The objective can be, for example, made by Zeiss of Jena, Germany, as well as other companies. The objective can be one of a plurality including 1X, 10X, 20X, 40X, and others, depending upon the application.
10 Magnification can be further expanded or contracted by intermediate optics between the objective and the camera. Selection of filter or filters is controlled by computer 51 via control line 75.

The camera 50 captures an image of cells from plate 59. The image is obtained from light scattering off of cells or portions of cells in the plate through
15 objective 53, through objective spacers, through filters 56, which are captured at camera 50. In this preferred embodiment, the camera is a digital camera, but can be an analogue camera. The digital camera is a CCD camera, which has 1280 by 1024 pixels, or more or less. The pixels can be 6.7 microns in dimension or more or less. The camera preferably is substantially free from an external shutter to quickly capture
20 a plurality of images of cells from the plate. The camera is controlled via control line 71 through controller 50A, which connects to computer 51 through control line 70. The present invention can also include other types of image acquisition devices selected from at least an epifluorescence, a confocal, a total-internal reflection, a phase, a Hoffman, a bright field, a dark field, a differential interference contrast, an
25 interference reflection, or multi-photon illumination device.

The present imaging system stores images on a high density memory device 60. The high density memory device is preferably optical, but can also be magnetic. The high density memory device can be any suitable unit that is capable of storing a plurality of images from a plurality of sites in the plate. The memory device
30 can be a compact disk, which would generally use a compact disk burner or the like. Depending upon the embodiment, the high density memory device is used to archive the images that are captured from the camera in the imaging system. Further details

of the imaging system can be found throughout the present specification, and more particularly below.

As merely an example, the present invention can be implemented using the following sequence of steps, which have been described in a journal entry form.

- 5 Here, images are opened and objects are identified based on a background value that has been edited in starting image acquisition. Information is maintained in a spreadsheet or other database format, which has the following information for each object:

Image Name	Image Plane	Image Date and Time
Elapsed Time	Object #	Total area
Pixel area	Area	Hole area
Relative hole area	Standard area count	Perimeter
Length	Breadth	Fiber length
Fiber breadth	Shape factor	Ell. form factor
Inner radius	Outer radius	Mean radius
Average gray value	Total gray value	Optical density
Radial dispersion	Texture Difference Moment	EFA Harmonic 2, Semi-Major Axis
EFA Harmonic 2, Semi-Minor Axis	EFA Harmonic 2, Semi-Major Axis Angle	EFA Harmonic 2, Ellipse Area
EFA Harmonic 2, Axial Ratio	EFA Harmonic 3, Semi-Minor Axis	

10

After computations are done, the log file is saved. In particular, the file is saved in an appropriate place with an appropriate name.

In a specific embodiment, the present invention provides the following detailed example of journal entries, which should not limit the scope of the invention.

Set Up Sequential File Names	Interactive: user sets up prefix name and image storage directory
Open Data Log	Opens a DDE (Excel) File
Annotate Log File	Interactive: experimental information that will go into the first line of the log file of stage positions
Stage (Go to Origin)	Origin is set as the center of well A1
Stage (Move to Absolute Position)	Offset to upper left hand corner of well (1410, 1621)
Stage (Log Position)	
Stage (Scan Wells)	User picks wells to scan: runs 3x3 image collection.jnl.

3X3 IMAGE COLLECTION.jnl

Stage (Scan)	Takes 9 images of well, -1600 motor steps apart from left to right 3 columns and 3 rows, runs FOCUS, COLLECT IMAGE, SAVE SEQUENTIAL FILE NAME.JNL.
--------------	--

5

FOCUS, COLLECT IMAGE, SAVE SEQUENTIAL FILE NAME.jnl

Stage (Log Position)	Logs stage position of each image
ADC – Focus	Opens up the manual focusing window with whatever focus time is current set
Show Message and Wait	Interactive: user hits enter to continue when done focusing

ADC-Acquire from Digital Camera	Takes Hoechst image
Save Using Sequential File Names	
Close	Closes image window

START IMAGE ANALYSIS.jnl

Low Pass	3x3 convolution of already opened image
Low Pass	3x3
Show Region Statistics	Interactive: Show entire image statistics. Calculate background subtraction value for step 4. by: INTENSITY Average + INTENSITY Std. Dev.
Arithmetic	Interactive: User inputs subtraction value from 3. into the constant Value field
Threshold image	Creates threshold 1 unit above 0 to 4096
Integrated Morphometry – Load State	Loads Start Image Analysis.ima Classifier 100 < area < 200000
Integrated Morphometry – Measure	Interactive: Shows area summary information about all objects. The average number is used as the Standard Area in 8.
Object Standards - Set Object Standards	Interactive: User inputs average area value from 7. into Standard Area box to be used by automated IMA for all images

IMA OBJECTS.jnl

Low Pass	3x3 convolution
Low Pass	3x3 convolution
Arithmetic	This background subtraction value needs to be manually entered into this journal from the value determined in START IMAGE ANALYSIS.jnl step 3
Threshold Image	1 unit above 0
Integrated Morphometry – Load State	Hoechst. IMA Classifier 200 < area < 200000
Integrated Morphometry – Measure	Measures statistical info for all objects
Run Journal	Runs log obj and sum data.jnl

Log obj and sum data.jnl

Integrated Morphometry – Log Data	Logs object data into Sheet 1
Integrated Morphometry – Log Data	Log summary data into Sheet 2

5

COLLECT AUTOMATED IMA DATA IN ONE SPREADSHEET.jnl

Run Journal	Runs OPEN OBJECT LOG DDE FILE.JNL
Loop for all Images in a Directory	Loops IMA OBJECTS.jnl
Close Summary Log	
Close Object Log	User must manually save Excel spreadsheet

OPEN OBJECT LOG DDE FILE.jnl

Open Object Log	Opens a DDE object log into sheet 1 of an Excel spreadsheet
Open Summary Log	Opens a summary log into sheet 2

COLLECT AUTOMATED IMA DATA IN ONE SPREADSHEET 16 BIT IMAGES.jnl

Arithmetic	Interactive: Opens Arithmetic window for user to input background subtraction level from START IMAGE ANALYSIS.jnl step 3
Run Journal	Runs OPEN OBJECT LOG DDE FILE.JNL
Loop for all Images in a Directory	Interactive: Runs IMA OBJECTS 16 bit.jnl. User picks directory from which to choose.

5

IMA OBJECTS 16bit.jnl

Low Pass	3x3 convolution
Low Pass	3x3 convolution
Copy to 8-bit Image	No autoscale, to new untitled image
Save Using Sequential File Name	Saves 8bit image using previously defined Sequential File names.
Arithmetic	This background subtraction value needs to be manually entered into this journal from the value determined in START IMAGE ANALYSIS16 TO 8 BIT.jnl step 5
Threshold Image	1 unit above 0 to 255

Integrated Morphometry – Load State	Hoechst.IMA Classifier 200 < area < 200000
Integrated Morphometry – Measure	Measures statistical info for all objects
Run Journal	Runs log obj and sum data.jnl

START IMAGE ANALYSIS 16 to 8 BIT.jnl

Copy to 8-bit Image	No autoscale, to new untitled image
Close	Closes 16 bit image
Low Pass	3x3 convolution
Low Pass	3x3 convolution
Show Region Statistics	Interactive: Show entire image statistics. Calculate background subtraction value for step 6. by: INTENSITY Average + INTENSITY Std. Dev.
Arithmetic	Interactive: User inputs subtraction value from 5. into the constant Value field
Threshold image	Creates threshold by 1 unit above 0 to 255
Integrated Morphometry – Load State	Loads Start Image Analysis.ima Classifier 100 < area < 200000
Integrated Morphometry – Measure	Interactive: Shows area summary information about all objects. The average number is used as the Standard Area in 10.
Object Standards - Set Object Standards	Interactive: User inputs average area value from 9. into Standard Area box to be used by automated IMA for all images

IMA OBJECTS WITH NEW LOG FILE.jnl

Run Journal	OPEN OBJECT LOG DDE FILE.JNL
Run Journal	IMA OBJECTS.jnl
Close Summary Log	
Close Object Log	User must manually save every Excel spreadsheet generated.

INTERACTIVE IMA OBJECTS.jnl

Threshold Image	User manually sets threshold
Integrated Morphometry – Load State	Hoechst.IMA Classifier $200 < \text{area} < 200000$
Integrated Morphometry – Measure	Objects
Integrated Morphometry – Log Data	Into open object.log file

5

COLLECT INTERACTIVE IMA DATA.jnl

Close Object Log	
Open Object Log	Interactive
Annotate Log File	Interactive: experimental information that will go into the first line of the object log file
Loop for all Images in Directory	Runs INTERACTIVE IMA OBJECTS.jnl

CHANGE FILTER, COLLECT IMAGE, SAVE SEQUENTIAL FILE
NAME.jnl

Stage (Log Position)	
ADC-Focus	

Show Message and Wait	Interactive – user presses Enter when done focusing
ADC – Acquire from Digital Camera	Hoechst
Save Using Sequential File Name	
Close	Close open image

COLLECT HOECHST AND FITC.jnl

Run Journal	FOCUS, COLLECT IMAGE, SAVE SEQUENTIAL FILE NAME.JNL
Run Journal	CHANGE FILTER, COLLECT IMAGE, SAVE SEQUENTIAL FILE NAME.jnl

3X3 IMAGE COLLECTION HOECHST FITC.jnl

Stage (Scan)	COLLECT HOECHST AND FITC.jnl
--------------	------------------------------

5

AUTOMATED 3X3 IMAGE COLLECTION HOECHST FITC.jnl

Set Up Sequential File Names	Interactive: user sets up prefix name and image storage directory
Open Data Log	Excel DDL files
Annotate Log File	Interactive: experimental information that will go into the first line of the log file of stage positions
Stage (Go to Origin)	Origin is set as the center of well A1
Stage (Move to Absolute Position)	Offset to upper left hand corner of well (1410, 1621)

Stage (Log Position)	
Stage (Scan Wells)	Interactive: user picks wells to scan: runs 3X3 IMAGE COLLECTION HOECHST FITC.jnl

AUTOMATED IMAGE COLLECTION.jnl

Set Up Sequential File Names	Interactive: user sets up prefix name and image storage directory
Open Data Log	Opens a DDE (Excel) File
Annotate Log File	Interactive: experimental information that will go into the first line of the log file of stage positions
Stage (Go to Origin)	Origin is set as the center of well A1
Stage (Log Position)	
Stage (Scan Wells)	Interactive: user picks wells to scan: runs FOCUS, COLLECT IMAGE, SAVE SEQUENTIAL FILE NAME.JNL. Well to well travel = (-9035, -9035)

5

STARTUP.jnl

Install and Configure Devices	Open Stage Meta Devices
Set Live Video Channel	

Preferences	<u>Measure Objects</u> : Draw failed classifier objects, Exclude objects that touch the edge of the image, Enable Elliptical Fourier Parameters, turn off Warn users when measurement data will be erased <u>Image Saving</u> : Save Tiff/stk using LZW compression <u>Image Windows</u> : Use transparent thresholds.
Configure Default Paths	C:\Metamorph Data C:\Metamorph Data\Common Settings
Load Journal Taskbar	Common.JTB

Nested Journals

Automated 3x3 Image Collection

- 5 *Loop* 3x3 image collection
 Loop focus, collect image, save sequential file name

Automated 3x3 image collection Hoechst FITC

- 10 *Loop* 3x3 image collection Hoechst FITC
 loop Collect Hoechst and FITC
 focus, collect image, save sequential file name
 change filter, collect image, save sequential file name

Automated image collection

- 15 *Loop* focus, collect image, save sequential file name

Collect automated IMA data in one Spreadsheet

Open object log DDE file

Loop IMA objects

Log obj and sum data

Collect automated IMA data in one spreadsheet 16 bit images

5 Open object log DDE file

Loop IMA objects 16 bit

Log obj and sum data

Although the above has been generally described in terms of a specific
10 user interface and software code, other user interfaces and code can also be used. One
of ordinary skill in the art would recognize many other variations, alternatives, and
modifications.

Fig. 6 is a simplified diagram 600 of a cleaning and dispensing system
according to an embodiment of the present invention. This system 600 includes a
15 variety of elements such as a dispensing head 609, which is coupled to a plurality of
pipettes 601. The pipettes input and output fluids or solutions from plate 603. The
plate has a plurality of sites, each of which can be used to input cells or a combination
of cells and solution. The system also has elements to house solutions 605, which are
used to manipulate cell samples in the plate. The dispensing head is supported
20 through a support member 607, which is sufficiently rigid to allow for movement of
the head. The dispenser is coupled to the present system in a mechanical and
electrical manner, which provides for a fully integrated system for providing cell
samples to the imaging system according to the present invention.

Fig. 7A illustrates a representative block flow diagram of simplified
25 process steps of a method for determining properties of a manipulation based upon
effects of the manipulation on one or more portions of one or more cells in a
particular embodiment according to the present invention. This diagram is merely an
illustration and should not limit the scope of the claims herein. One of ordinary skill
in the art would recognize other variations, modifications, and alternatives. In step
30 700, one or more samples of cells can be provided. These cells can be live, dead, or
fixed cells, or cell fractions. The cells also can be in one of many cell cycle stages,
including G0, G1, S, G2 or M phase, M phase including the following cell cycle
stages: interphase, prophase, prometaphase, metaphase, anaphase, and telophase.

Cell components tracked in presently preferable embodiments can include proteins, protein modifications, genetically manipulated proteins, exogenous proteins, enzymatic activities, nucleic acids, lipids, carbohydrates, organic and inorganic ion concentrations, sub-cellular structures, organelles, plasma membrane, adhesion complex, ion channels, ion pumps, integral membrane proteins, cell surface receptors, G-protein coupled receptors, tyrosine kinase receptors, nuclear membrane receptors, ECM binding complexes, endocytotic machinery, exocytotic machinery, lysosomes, peroxisomes, vacuoles, mitochondria, Golgi apparatus, cytoskeletal filament network, endoplasmic reticulum, nuclear membrane, proteosome apparatus, chromatin, nucleolus, cytoplasm, cytoplasmic signaling apparatus, microbe specializations and plant specializations.

The following table illustrates some markers and cell components commonly used by embodiments according to the present invention. Other markers can be used in various embodiments without departing from the scope of the invention.

Cell component	Marker	Disease State
Plasma membrane (including overall cell shape)	Carbocyanine dyes Phosphatidylserine Various lipids Glycoproteins	Apoptosis-Cancer Apoptosis-Neural degenerative Ds
Adhesion complexes	Cadherins Integrins Occludin Gap junction ERM proteins CAMs Catenins Desmosomes	Thrombosis Metastasis Wound healing Inflammatory Ds Dermatologic Ds
Ion Channels and Pumps	Na/K Atpase Calcium channels Serotonin reuptake pump CFTR	Cystic fibrosis Depression Congestive Heart Failure Epilepsy

G coupled receptors	β adrenergic receptor Angiotensin receptor	Hypertension Heart Failure Angina
Tyrosine kinase receptors	PDGF receptor FGF receptor IGF receptor	Cancer Wound healing Angiogenesis Cerebrovascular Ds
ECM binding complexes	Dystroglycan Syndecan	Muscular Dystrophy
Endocytotic machinery	Clathrin Adaptor proteins COPs Presenilins Dynamin	Alzheimer's Ds
Exocytotic machinery	SNAREs Vesicles	Epilepsy Tetanus Systemic Inflammation Allergic Reactions
Lysosomes	Acid phosphatase Transferrin	Viral diseases
Peroxisomes/Vacuoles		Neural degenerative Ds
Mitochondria	Caspases Apoptosis inducing factor F1 ATPase Fluorescein Cyclo-oxygenase	Apoptosis Neural degenerative Ds Mitochondrial Cytopathies Inflammatory Ds
Golgi Apparatus	Lens Culinaris DiOC6 carbocyanine dye COPs	

Cytoskeletal Filament Networks	Microtubules	Cancer
	Actin Intermediate Filaments Kinesin, dynein, myosin Microtubule associated proteins Actin binding proteins Rac/Rho Keratins	Neural degenerative Ds Inflammatory Ds Cardiovascular Ds Skin Ds
Endoplasmic Reticulum	SNARE PDI Ribosomes	Neural degenerative Ds
Nuclear Membrane	Lamins Nuclear Pore Complex	Cancer
Proteosome Apparatus	Ubiquityl transferases	Cancer
Chromatin	DNA Histone proteins Histone deacetylases Telomerases	Cancer Aging
Nucleolus	Phase markers	
Cytoplasm	Intermediary Metabolic Enzymes BRCA1	Cancer
Cytoplasmic Signaling Apparatus	Calcium Camp PKC pH	Cardiovascular Ds Migraine Apoptosis Cancer
Microbe Specializations	Flagella Cilia Cell Wall components: Chitin synthase	Infectious Ds

Plant specializations	Choloroplast	Crop Protection
	Cell Wall components	

Then, in a step 702, one or more samples of the manipulation can be provided to the cells. Manipulations can comprise one or any combination of chemical, biological, mechanical, thermal, electromagnetic, gravitational, nuclear, or temporal factors, for example. For example, manipulations could include exposure to chemical compounds, including compounds of known biological activity such as therapeutics or drugs, or also compounds of unknown biological activity. Or exposure to biologics that may or may not be used as drugs such as hormones, growth factors, antibodies, or extracellular matrix components. Or exposure to biologics such as infective materials such as viruses that may be naturally occurring viruses or viruses engineered to express exogenous genes at various levels. Bioengineered viruses are one example of manipulations via gene transfer. Other means of gene transfer are well known in the art and include but are not limited to electroporation, calcium phosphate precipitation, and lipid-based transfection. Manipulations could also include delivery of antisense polynucleotides by similar means as gene transfection. Other genetic manipulations include gene knock-outs or gene mutations. Manipulations also could include cell fusion. Physical manipulations could include exposing cells to shear stress under different rates of fluid flow, exposure of cells to different temperatures, exposure of cells to vacuum or positive pressure, or exposure of cells to sonication. Manipulations could also include applying centrifugal force. Manipulations could also include changes in gravitational force, including sub-gravitation (the preferred embodiment in outer space). Manipulations could include application of a constant or pulsed electrical current. Manipulations could also include irradiation. Manipulations could also include photobleaching which in some embodiments may include prior addition of a substance that would specifically mark areas to be photobleached by subsequent light exposure. In addition, these types of manipulations may be varied as to time of exposure, or cells could be subjected to multiple manipulations in various combinations and orders of addition. Of course, the type of manipulation used depends upon the application.

Then, in a step 704, one or more descriptors of a state in the portions of the cells in the presence of the manipulation can be determined using the images

collected on the imaging system. Descriptors can comprise scalar or vector values, representing quantities such as area, perimeter, dimensions, intensity, gray level, aspect ratios, and the like. Other types of descriptors include, but are not limited to, one or any combination of characteristics such as a cell count, an area, a perimeter, a
 5 length, a breadth, a fiber length, a fiber breadth, a shape factor, a elliptical form factor, an inner radius, an outer radius, a mean radius, an equivalent radius, an equivalent sphere volume, an equivalent prolate volume, an equivalent oblate volume, an equivalent sphere surface area, an average intensity, a total intensity, and an optical
 10 density. These descriptors can be average or standard deviation values, or frequency statistics from the descriptors collected across a population of cells. These descriptors can be further reduced using other methods such as principal component analysis and the like. In some embodiments, the descriptors include features from different cell portions or cell types. That is, a first feature can be from a nuclei and a second feature is from another cell structure such as Golgi apparatus, mitochondria, spacing between
 15 cell structures or cells themselves, as well as many others.

A presently preferable embodiment uses descriptors selected from the following table. Other descriptors can also be used without departing from the scope of the invention.

Name of Parameter	Explanation/Comments
Count	Number of objects
Area	
Perimeter	
Length	X axis
Width	Y axis
Shape Factor	Measure of roundness of an object
Height	Z axis
Radius	
Distribution of Brightness	
Radius of Dispersion	Measure of how dispersed the marker is from its centroid
Centroid location	x-y position of center of mass
Number of holes in closed objects	Derivatives of this measurement might include, for

	example, Euler number (= number of objects - number of holes)
Elliptical Fourier Analysis (EFA)	Multiple frequencies that describe the shape of a closed object
Wavelet Analysis	As in EFA, but using wavelet transform
Interobject Orientation	Polar Coordinate analysis of relative location
Distribution Interobject Distances	Including statistical characteristics
Spectral Output	Measures the wavelength spectrum of the reporter dye. Includes FRET
Optical density	Absorbance of light
Phase density	Phase shifting of light
Reflection interference	Measure of the distance of the cell membrane from the surface of the substrate
1,2 and 3 dimensional Fourier Analysis	Spatial frequency analysis of non closed objects
1,2 and 3 dimensional Wavelet Analysis	Spatial frequency analysis of non closed objects
Eccentricity	The eccentricity of the ellipse that has the same second moments as the region. A measure of object elongation.
Long axis/Short Axis Length	Another measure of object elongation.
Convex perimeter	Perimeter of the smallest convex polygon surrounding an object
Convex area	Area of the smallest convex polygon surrounding an object
Solidity	Ratio of polygon bounding box area to object area.
Extent	proportion of pixels in the bounding box that are also in the region
Granularity	
Pattern matching	Significance of similarity to reference pattern
Volume measurements	As above, but adding a z axis

Then, in a step 705, a database of cell information can be provided.

Next, in a step 706, a plurality of descriptors can be searched from a database of cell information in order to locate descriptors based upon one of the descriptors of the manipulation. Then, in a step 708, properties of the manipulation are predicted based upon the properties of the located descriptors. Properties can comprise toxicity, specificity against a subset of tumors, mechanisms of chemical activity, mechanisms of biological activity, structure, adverse biological effects, biological pathways, clinical effects, cellular availability, pharmacological availability, pharmacodynamic properties, clinical uses and indications, pharmacological properties, such as absorption, excretion, distribution, metabolism and the like.

In a particular embodiment, step 706 comprises determining matching descriptors in the database corresponding to a prior administration of the manipulation to the descriptors of the present administration of the manipulation. In a particular embodiment according to the present invention, combinations of measurements of scalar values can provide predictive information. A database can be provided having one or more "cellular fingerprints" comprised of descriptors of cell-substance interactions of drugs having known mechanisms of action with cells. Such descriptors can be analyzed, classified, and compared using a plurality of techniques, such as statistical classification and clustering, heuristic classification techniques, a technique of creating "phylogenetic trees" based on various distance measures between descriptors from various drugs. In this embodiment, numeric values for the descriptors can be used by comparison techniques. A phylogenetic tree can be created that illustrates a statistical significance of the similarity between descriptors for the drugs in the database. Because the drugs used to build the initial database are of known mechanism, it can be determined whether a particular scalar value in a descriptor is statistically predictive. Finally, a compound descriptor with no known mechanism of action can be queried against the database and be statistically compared and classified among the drugs in the database that the compound most resembles.

In a particular embodiment, relationships between measured morphological properties of images and physiological conditions can be determined. Relationships can include, for example, treatment of different cell lines with chemical compounds, or comparing cells from a patient with control cells, and the like. In a presently preferable embodiment, comparisons can be performed on acquired image

features. Some embodiments can comprise statistical and neural network - based approaches to perform comparisons of various features. The foregoing is provided as merely an example, and is not intended to limit the scope of the present invention. Other techniques can be included for different types of data.

5 In some embodiments, classification, clustering and other types of predictive data analysis can be performed on features extracted from cell images. In a presently preferable embodiment, statistical procedures for comparisons, classification and clustering are performed on data obtained from imaging cells.

Fragments of data preparation and pre-formatting (S language):

```
10 >tmp.frame <- Generic.Summary  
>names1 <- paste("Cell.line.5", tmp.names, sep=".")  
> by.compound.matrix <- as.matrix(arranged.by.compound)
```

Example of the code for principal component analysis (data
15 preparation) using S language:

```
all.data.princomp <- menuPrincomp(data =  
by.compound.matrix, scores = T, cor = "Correlation",  
na.action = T, print.short = T, print.importance = T,  
print.loadings = T, cutoff.loadings = 0.1,  
20 plot.screepplot = T, plot.loadings = T, plot.biplot = T,  
plot.biplot.choices = c(1,2), predict.p = F)
```

Example of clustering using a divisive hierarchical clustering
algorithm:

```
25 > div.hier.2.manhattan.cluster$call  
diana(x = tmp.sum.by.comp, diss = F, metric =  
"manhattan",  
stand = T, save.x = T, save.diss = T)
```

30 Another embodiment utilizes existing tools for biological sequence similarity searches, classification, and phylogenetic analysis. In a particular embodiment, numbers in a numerical descriptor can be substituted by one or more of nucleic acid or amino acid codes according to a one of several sets of rules. Once

converted into a corresponding nucleotide or amino acid sequence representation, the fingerprints can be analyzed and compared using software and algorithms known in the art for genetic and peptide sequence comparisons, such as GCG, a product of Genetics Computer Group, with company headquarters in Madison WI. Select
5 embodiments comprising such approaches enable the use of a broad array of sophisticated algorithms to compare, analyze, and cluster gene and protein sequences. Many programs performing this task are known to those of ordinary skill in the art, such as for example, the PHYLIP (PHYlogeny Interference Package) a package of programs for inferring phylogenics (evolutionary trees) described in (Feldenstein, J.
10 1996 Methods Enzymol 266:418-427 and Feldenstein, J. 1981 J. Mol. Evol. 17(6):368-376).

Embodiments can perform such analysis based upon factors such as numerical value, statistical properties, relationships with other values, and the like. Further details of a step of manipulation are noted more particular below.

15 Fig. 7B illustrates a representative block flow diagram of simplified process steps for determining one or more descriptors of a state in the portions of the cells in the presence of the manipulation of step 704 of Fig. 7A in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill
20 in the art would recognize other variations, modifications, and alternatives. In a step 712, an image of a cell portion is obtained. In some embodiments, the cell portion is visualized with a fluorescently labeled marker that is specific for the portion or portions of interest. A cell portion can include, for example, one or more of the following: nuclei, Golgi apparatus, and other features. The cell portion may vary in
25 select embodiments according to the invention. Then, in a step 714, a digitized representation of the image obtained in step 712 is determined. In some embodiments, steps 714 and step 712 can comprise a single step. These embodiments use a digital imaging means such as a digital camera, to obtain a digital image of the target directly. Next, in a step 716, the digital representation of the image is
30 processed to obtain image features. Image features can include such quantities as area, perimeter, dimensions, intensity, aspect ratios, and the like. Then, in a step 718 descriptors can be determined from the image features. Descriptors can comprise scalar or vector quantities and can comprise the image features themselves, as well as

composed features, such as shape factor derived by a relationship $4\pi * \text{area} / \text{perimeter}$, and the like. Descriptors can also comprise statistical quantities relating to feature characteristics across a population of cells, such as a standard deviation, and average, and the like.

5 In a preferred embodiment, cells can be placed onto a microscope, such as a Zeiss microscope, or its equivalent as known in the art. A starting point, named Site A01, is identified to the microscope. A plurality of exposure parameters can be optimized for automated image collection and analysis. The microscope can automatically move to a new well, automatically focus, collect one or more images, at
10 one or more wavelengths, move to a next well, and repeat this process for all designated wells in a multiple well plate and for multiple plates. A file having a size and an intensity distribution measurement for each color and rank for each well can then be created for the images acquired. Based on this information, a user or a computer can revisit sites of interest to collect more data, if desired, or to verify
15 automated analysis. In a presently preferred embodiment, image automatic focus and acquisition can be done using computer software controlling the internal Z-motor of the microscope. Images are taken using a 10x, 20x, or 40x air long working distance objectives. Sometimes multiple images are collected per well. Image exposure times can be optimized for each fluorescent marker and cell line. The same exposure time
20 can be used for each cell line and fluorescent marker to acquire data.

Fig. 7C illustrates a representative block flow diagram of simplified process steps for obtaining images of cell portions of step 712 of Fig. 7B in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill
25 in the art would recognize other variations, modifications, and alternatives. The method is generally outlined by the steps below:

- (1). In a step 720, a sample is provided to the imaging device. Samples can be provided in 96 well plates and the like. The sample may be loaded into a microscope, such as a Zeiss microscope or equivalent.
- 30 (2). In a step 722, a set of optical filters is selected to shine light of the appropriate wavelength to illuminate the first sample, which may be contained in a first well designated A01.

(3). In a step 724, an automatic focusing procedure is performed for the site. In a particular embodiment, the internal z-motor of the microscope which is attached to the objective nosepiece is used for automatic focusing of the microscope. In an alternative embodiment, the plate holding the samples is moved to perform automatic focusing of the microscope, or focusing can be performed by moving optical components attached to the microscope and the like.

(4). In a step 726, images are collected for the site. Images can be collected for every color at every site. Present embodiments can provide images for up to four colors. However, embodiments are contemplated that can provide more colors by using either a monochromator coupled with excitation filters which are on a filter wheel, or by digitally separating overlapping fluorophores. Those knowledgeable in the field will know that given calibration images of single fluorophores, a look-up table can be devised which will allow for the digital removal of fluorescence bleed-through of fluorescence which may occur in optical channels other than the one for which that filter has been optimized in instances of using more than one fluorophore at once. Cell growth and density information is also collected. Cell density is determined by what percentage of the area being imaged is inhabited by cells. In some embodiments, imaging can be facilitated using one or more biosensors, molecules such as non-proteins, i.e., lipids and the like, that are luminescently tagged. However, some embodiments can also use fluorescence polarization and the like. Fluorescence polarization is a homogeneous fluorescence technology where the excited state of the molecule lasts much longer than in normal fluorescence, taking seconds to minutes to reach equilibrium, obliterating the need to wash away fluorescence markers that are not specifically bound to a marker. Further, embodiments can detect differences in spectral shifts of luminescent markers. Some fluorescence markers, such as Nile Red sold by Molecular Probes of Eugene, OR, will change its emission peak wavelength depending on its environment. One can detect these changes by monitoring the level of fluorescence at both wavelengths and reading out at ratio of the two.

(5). In a step 728, a determination is made whether more fields of view need to be taken for a particular color. If this is so, then processing continues at step 726 at a new site. Otherwise, processing continues with a decisional step 730.

Images can now be taken by repeating step 726. In a preferred embodiment 4 to 9 images are collected at each site.

(5). In a step 730, a determination is made whether more optical configurations need to be taken in order to obtain images for all differently-marked cell portions the sample. If this is so, then in a step 732 a new optical configuration is determined. Images for the new optical configuration can now be taken by repeating steps 726 and 728.

(6). In a decisional step 734, after all optical configurations and images for fields of view in a sample have been obtained, a determination is made whether any further samples remain to be analyzed. If so, a new sample is brought into view and processing continues with step 720. Otherwise, image processing is complete. In a presently preferable embodiment, image data can be stored on a CD ROM using a CD ROM burner, such as CRW4416 made by Yamaha of Japan. However, other mass storage media can also be used.

Fig. 7D illustrates a representative block flow diagram of simplified process steps for processing digitized representations of step 716 of Fig. 7B in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. The method is generally outlined by the steps below:

(1). In a step 740, a digitized image input is preprocessed. Preprocessing might include, but is not limited to, such operations as background subtraction, thresholding, smoothing, adoptive filtering, edge enhancements, contrast enhancements, histogram equalization. A particular combination of preprocessing steps can be applied to images in successive steps or in parallel to copies of the image.

A simplified example of a smoothing and background subtraction procedure in a MatLab language is presented in computer code below:

```
function Isubtracted = cmBackgrSubtrl(I,k)
% cmBackgrSubtrl(I,k) - simple flat background (=modal*k)
subtraction
% Y = cmBackgrSubtrl(I, k) - image Y is generated by
```



```

    % subtraction (with saturation) of modal pixel value of I
    multiplied by k
    % DEFAULT - k=1
    %
5   if (nargin == 1)
        k=1;
    end
    if (size(k)~=1)
        error('cmBackgrSubtrl: parameter k should be a number.
10   Exiting...');
    end

    %modpixnum = floor(size(I(:),1)/2);
    %sortedval = sort( double(I(:)) );
15  %modpixel = sortedval(modpixnum);
    modpixel = median(double(I(:)));
    bg = k*modpixel;

    Isubtracted = mmsubm( uint8(I), uint8(round(ones(
20  size(I))*k*modpixel )) );

```

An example of a procedure for thresholding in computer code (MatLab) is presented below:

```

function thresh = GetThreshByPerim1(I, M)
25  % GetThreshByPerim1(I) Finds optimal thresholding value
    for image I
    % N = GetThreshByPerim1(I) Finds thresholding value N for
    image I
    % N = GetThreshByPerim1(I, M) - tests threshold values up
30  to M
    % DEFAULT M = maximum pixel value in I
    % note that GetThreshByArea is significantly faster
    % finds a threshold value that causes the maximal change
    in the

```

```
% total perimeter of the objects (Russ ????)
% see Matlab_Auto_threshold1_1-23-99.doc for more details
% Note: works somewhat better on SMOOTH images (i.e.
medfilt2(I, [3 3]) two times

5
if (nargin == 0)
    error (strcat( mfilename, ' : at least one parameter
required')));
elseif (nargin == 1)
10    M = double(max(I(:)));      %test thresholds up to
    maximum pixel value in I
    elseif (nargin > 2)
        error (strcat (mfilename, ' : too many parameters'));
    end

15
if (size(M)>1)
    error (strcat(mfilename, ' : argument M should be a
number')));
end

20
Minval = double( min(I(:)));
step = 1;

%generate vertical vector perims with total perimeters of
25 objects at different
%threshold values
for i=Minval : step : M
    bwI = im2bw(I, i/255);
    prI = bwperim(bwI);
30    pr = sum(prI(:));
    if (exist('perims', 'var') == 0) %perims is yet
undefined
        perims = pr;
    else
```

```

        perims = cat(1, perims, pr);
    end
end

5  % vector prdiffs contains differences between successive
    perimeters
    prdiffs = diff(perims);
    mindecrease = min(prdiffs);
    minvalues = find(prdiffs == mindecrease);
10  index_of_mindecrease = minvalues(1);
    thresh = index_of_mindecrease + 1;

    % =====end GetThresh1=====

```

15 Thresholding provides a specific intensity, such that pixels darker than the threshold are deemed black, and pixels lighter than the threshold are considered white. The thresholded image can be processed using binary image processing techniques in order to extract regions.

(2). In a step 742+744, the digitized image input is subjected to object
20 identification. This can be accomplished by a variety of procedures, for example by thresholding or edge detection and subsequent morphological opening and closing. Edge detection can be accomplished by means of gradient-based or zero-crossing methods, such as Sobel, Canny, Laplassian, Perwitt, and other methods.

An example of object identification procedure based on Canny edge
25 detection (in MatLab language) is presented below:

```

function Imask = cmMaskDNA1(I);
% cmMaskDNA1 - generates binary mask for cell nuclei
through edge detection
30 % Imask = cmMaskDNA1(I)
% PARAMETERS
%   I - intensity image (grayscale)
% OUTPUT
%   Imask - BW image with objects from I

```

```

%
% For more details see Notebook Matlab_DNA_masking1_1-22-
99.doc
% Uses SDC Morphology Toolbox V0.7

5
if (nargin ~= 1)
    error('Wrong number of input parameters');
end
if (nargout ~= 1)
10    error('Wrong number of output parameters: one output
argument should be provided');
end

15    Imask = edge(I, 'canny');
    Imask = mm dil(Imask, mmsecross(1));
    Imask = mmero ( mmc lohole(Imask, mmsecross(1)));
    Imask = mmedgeoff(Imask, mmsecross(1));
    % note that mmedgeoff this command removed FILLED OBJECTS
20    but not touching OUTLINES.
    % these outlines can be removed by filtering:
    Imask = medfilt2(Imask, [5 5]);

    %=====end cmMaskDNA1
25    =====

```

However, embodiments can also use other techniques, such as Fast Fourier Transforms (FFT) and the like as known in the art without departing from the scope of the present invention.

30 (3). In a step 746, a plurality of region features can be determined. For example, in a representative embodiment, image features can include such quantities as area, perimeter, dimensions, intensity, aspect ratios, and the like. Features not directly related to individual objects are also being extracted.

An example of a procedure for extraction of some of the features (MatLab language) is presented below:

```

function OData = cmGetObjectsData(I, Ilabel)
5  % cmGetObjectsData returns array measurements of objects
  in image "I" masked by "Ilabel"
  % EV 2-3-99; 2-10-99
  % OData = cmGetObjectsData(I, Ilabel) returns an array of
  morphological and intensity measurements
10  %   taken from a grayscale image "I". Objects are
  identified on a mask image Ilabel, usually
  %   created by bwlabel()
  % OUTPUT:
  % Each row in the output array OData represents
15  individual object
  % columns contain the following measurements:
  %
  %   1 - Index ("number" of an object);      8 -
  Solidity;
20  %   2 - X coordinate of the center of mass; 9 - Extent;
  %   3 - Y coordinate      "-"; 10 - Total
  Intensity;
  %   4 - Total Area (in pixels);              11 - Avg.
  Intensity;
25  %   5 - Ratio of MajorAxis/MinorAxis;      12 - Median
  Intensity;
  %   6 - Eccentricity;                        13 - Intensity of
  20% bright pixel
  %   7 - EquivDiameter;                       14 - Intensity of
30  80% bright pixel
  %
  % For details on morphological parameters see information
  on MatLab imfeature();

```

```
% Intensity parameters are either obvious or are
documented in comments in this file.

if (nargin ~= 2)
5   error ('function requires exactly 2 parameters');
end
if (nargout ~= 1)
    error ('function has 1 output argument (array X by
14) ');
10 end

% finished checking arguments

% first collect morphological parameters in a structure
15 array:
ImStats = imfeature(Ilabel, 'Area', 'Centroid',
    'MajorAxisLength',...
        'MinorAxisLength', 'Eccentricity', 'EquivDiameter',
    ...
20   'Solidity', 'Extent', 8 );

% now convert it into array (matrix) while collecting
intensity data for each object:

25 %preallocate output array:
numobjects = size(ImStats, 1);
OData = zeros(numobjects, 14);
%now convert ImStats into array and add intensity data to
it
30 for k=1:numobjects
    OData(k, 1) = k;
    OData(k, 2) = ImStats(k).Centroid(1);
    OData(k, 3) = ImStats(k).Centroid(2);
    OData(k, 4) = ImStats(k).Area;
```

```

        OData(k, 5) = (ImStats(k).MajorAxisLength) /
        (ImStats(k).MinorAxisLength);
        OData(k, 6) = ImStats(k).Eccentricity ;
        OData(k, 7) = ImStats(k).EquivDiameter;
5       OData(k, 8) = ImStats(k).Solidity;
        OData(k, 9) = ImStats(k).Extent;

        % now collect and assign intensity parameters from
        image I
10
        object_pixels = find( Ilabel == k);
        object_area = size(object_pixels, 1); %same as total
        number of pixels in the object
        object_intensities = double(I(object_pixels)); %
15 need to convert to double to do math
        sorted_intensities = sort(object_intensities); %
        will need to get median, 20% and 80% pixels
        total_intensity = sum(object_intensities, 1);
        avg_intensity = total_intensity / object_area;
20 median_intensity = sorted_intensities( floor(
        object_area/2 ) + 1 );
        pix20 = sorted_intensities( floor(object_area*0.2)+1
        ) ; %brightest pixel among dimmest 20%
        pix80 = sorted_intensities( floor(object_area*0.8)+1
25 ) ;

        OData(k, 10) = total_intensity;
        OData(k, 11) = avg_intensity;
        OData(k, 12) = median_intensity;
30 OData(k, 13) = pix20; %brightest pixel among dimmest
        20%
        OData(k, 14) = pix80; %dimmest pixel among brightest
        20%
        end %for

```

```
%===== end function  
cmGetObjectsData() =====
```

5 (4). In a step 748, quantitative descriptors characterizing cell state are calculated based on the feature measurements extracted at step 746. For example, histogram distribution of intensities of cell nuclei provides information about the population cell cycle stages.

10 In a particular embodiment according to the present invention, data analysis techniques for describing the fluorescence patterns of cell portions in multiple cell lines in the presence and absence of compounds are provided. Automated image analysis techniques can include determining one or more regions from around nuclei, individual cells, organelles, and the like, called "objects" using a thresholding function. Objects that reside on the edge of an image can be included or
15 excluded in various embodiments. An average population information about an object can be determined and recorded into a database, which can comprise a database text file or Excel spreadsheet, for example. However, embodiments can use any recording means without departing from the scope of the present invention. Values measured can be compared to the visual image. One or more types of numerical
20 descriptors can be generated from the values. For example, descriptors such as a number of objects, an average, a standard deviation of objects, a histogram (number or percentage of objects per bin, average, standard deviation), and the like can be determined.

25 In a particular embodiment according to the present invention, data can be analyzed using morphometric values derived from any of a plurality of techniques commonly known in the art. For example, a software package called MetaMorph Imaging System, provided by Universal Imaging Corporation, a company with headquarters in West Chester, PA and NIH Image, provided by Scion Corporation, a company with headquarters in Frederick, Maryland.

30 Fluorescent images can be described by numerical values, such as for example, an area, a fluorescence intensity, a population count, a radial dispersion, a perimeter, a length, and the like. Further, other values can be derived from such measurements. For example, a shape factor can be derived according to a relationship

$4\pi \cdot \text{area} / \text{perimeter}$. Other values can be used in various embodiments according to the present invention. Such values can be analyzed as average values and frequency distributions from a population of individual cells.

In a particular embodiment according to the present invention, techniques for the automatic identification of mitotic cells are provided. Image analysis techniques employing techniques such as multidimensional representations, frequency-based representations, multidimensional cluster analysis techniques and the like can be included in various embodiments without departing from the scope of the present invention. Techniques for performing such analyses are known in the art and include those embodied in MatLab software, produced by MathWorks, a company with headquarters in Natick, MA.

Scalar values providing efficacious descriptors of cell images can be identified using the techniques of the present invention to perform predictive analysis of drug behavior. In a presently preferred embodiment, a plurality of heterogeneous scalar values can be combined to provide descriptors for each manipulation. By applying predictive analysis routines to the collections of these descriptors, predictive information about any number of manipulations and cell interactions can be extracted.

Fig. 7E illustrates a representative block flow diagram of simplified process steps for analyzing image feature values to obtain descriptors of cell state of step 718 of Fig. 7B in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. Fig. 7E illustrates an input data of descriptors of known manipulations 319. A step 320 of reformatting and transforming data 319 to formats suitable for analysis is performed. Additionally, a "cleaning" process can eliminate outlying data points and the like in the data. Then, in a step 322, a decision is made whether to continue with step 324 or with step 326 based upon determining a particular type of analysis appropriate for the present application or particular type of prediction. If decisional step 322 determines processing should continue with step 324, then, in that step, an error estimate using a set of test descriptors is performed to estimate the quality of a prediction and processing continues with step 320. Once an optimal prediction is achieved, processing continues with step 326. In step 326, optimal transformation parameters and prediction methods are selected for use in

steps 328 and 330 which analyze data about an unknown manipulation. In a step 328, a solution is generated based upon any of techniques including training a neural network, solving a mathematical equation, applying decision tree rules and/or the like. In a step 330, an input data set of unknown descriptors 318 is reformatted and

5 transformed based upon the optimal transformation parameters selected in step 326 using the transformation procedures in steps 320, 322 and 324. In a step 332, predictions techniques are applied to the reformatted manipulations from step 330 and the solution generated in step 328 and a plurality of properties of known manipulations 317 (e.g., therapeutic properties, and the like) in order to determine a

10 prediction of properties of unknown manipulation 316.

Fig. 7F illustrates a representative block flow diagram of simplified process steps for a method of mapping a manipulation of cells to a physiological characteristic in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein.

15 One of ordinary skill in the art would recognize other variations, modifications, and alternatives. The method is generally outlined by the steps below:

(1) In a step 750, a plurality of cells, e.g., dead, live, cell fractions or mixtures of cells are provided.

(2) Then, in a step 752, the plurality of cells is manipulated, where

20 manipulation occurs using a source(s) from one or a combination selected from an electromagnetic, electrical, chemical, thermal, gravitational, nuclear, temporal, or a biological source.

(3) Next, in a step 754, a feature value is captured from the plurality of cells. The feature value can include one or any combination of characteristics such as

25 cell count, area, perimeter, length, breadth, fiber length, fiber breadth, shape factor, elliptical form factor, inner radius, outer radius, mean radius, equivalent radius, equivalent sphere volume, equivalent prolate volume, equivalent oblate volume, equivalent sphere surface area, average intensity, total intensity, and optical density. This list is not meant to be limiting.

(4) Then, in a step 756, a degree of presence of one or more feature values is assigned for each manipulation.

30

(5) In a step 758, the feature values from the plurality of cells are stored in memory locations. From the memory locations the values can be used for

statistical analyses to produce predictive information about the relatedness of the descriptors of the manipulations to one another. This information is used to infer properties of the manipulations.

Fig. 7G illustrates a representative block flow diagram of a simplified process steps for a method for populating a database with manipulated biological cell information in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. The method is generally outlined by the steps below:

10 (1) In a step 760, a plurality of cells in various stages of the cell cycle, A montage image that was used as a source to generate data in Appendix A is presented in Fig. 12., such as for example, the stages of interphase, prophase, metaphase, anaphase, and telophase are provided.

(2) Then, in a step 762, each of the cells in the various stages of mitotic development is manipulated.

(3) Next, in a step 764, an image of the plurality of manipulated cells is captured using image acquisition techniques in order to provide a morphometric characteristic of each of the manipulated cells.

20 (4) As a preferable option, in a step 766, an image database may be populated with the image of the plurality of manipulated cells.

(5) Following step 764 or optional step 766, a morphological value is calculated from the image in a step 768.

(6) In a step 770, the database is populated with the morphological value.

25 Fig. 7H illustrates a representative block flow diagram of simplified process steps for a method for populating a database with manipulated biological information, e.g., image acquisition parameters, image feature summary information, and well experimental parameters in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. Fig. 7H illustrates a step 780 in which cells are placed into site on a plate and a manipulation is applied. Then, in a step 781 an image is taken of the cells. In step 782, the image is transferred to an image archive

30

database. Then, in a step 783, well experimental parameters are entered into the database 787. Well experimental parameters can include cell type, manipulation and the like. In a step 784, image acquisition parameters are transferred to database 787. Image acquisition parameters can include file name, fluorophores and the like. In a
5 step 785, the image acquired in step 781 is analyzed. Then, in step 786, an image feature summary from the analysis step 785 is transferred to database 787.

In step 788, a lookup table for all analyses is provided to database 787. The lookup table provides information about the analyses. In a step 789, a query of database 787 for process data is performed. The results are reformatted. Then in a
10 step 790, the database 787 is queried. Next, in a step 791, features of the manipulations stored in the database are combined and reduced. Next, in a step 793, reduced features of step 791 can be compared. In a step 792, the results of step 793 are recorded in database 787. Then, in a step 794, a report of predictions based on comparisons performed in step 793 is generated.

15 Fig. 7I illustrates a representative block flow diagram of simplified process steps for acquiring images of manipulated biological information, e.g., cells, cell tissues, and cell substituents in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations,
20 modifications, and alternatives. Fig. 7I illustrates a step 770 in which a user sets up an image analysis procedure. Then, in a step 772, an image is read into image analysis software. Next, in a step 774, patterns and objects are identified in the image using one or more algorithms. Next, in a step 776, sets of features are extracted from the image. Then, in a step 778, feature information, descriptor values and the like are
25 exported to the database, such as database 787 of Fig. 7H, for recording. Next, in a decisional step 779, a determination is made whether any more images should be taken. If this is so, processing continues with step 772. Otherwise, image acquisition processing is completed.

Fig. 7J illustrates a representative block flow diagram of simplified
30 process steps for populating, acquiring and analyzing images of manipulated biological information in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations,

modifications, and alternatives. Fig. 7J illustrates a step 300 of placing a plate onto an imaging stage and reading a bar code. Then, in a step 301 an autofocus procedure is performed. Next, in a step 302, a first optical filter configuration is selected and an image is collected. Then, in a decisional step 303, a determination is made whether
5 more than one image per optical configuration can be taken. If so, then, in a step 304, a new position within the well is targeted and another image is collected. Then, in a decisional step 305, a determination is made whether any more images need to be collected. If this is so, step 304 is repeated until all images for a particular well have been collected. After one or more images are collected for the well, in a step 306, the
10 stage is returned to a starting position within the well, and a montage is created from collected images. The results are named with a unique file name and stored.

In a decisional step 307, a determination is made whether any more optical channels in the well can be imaged. If this is so, then in a step 308 the next optical filter configuration is selected and an image is collected. Processing then
15 continues with decisional step 303, as described above. Otherwise, if no further optical channels in the well can be imaged, then in a decisional step 309 a determination is made whether any wells remain to be imaged. If not all wells have been imaged, then in a step 310, the stage moves to the next well and processing continues with step 301, as described above. Otherwise, if all wells on the plate have
20 been imaged, then in a decisional step 311, a determination is made whether any more plates can be processed. If this is so, then processing continues with step 300 as described above. Otherwise, in a step 312, the information is stored to a CD or other storage device as a backup.

Fig. 7K illustrates a representative block flow diagram of simplified
25 process steps compound based upon information about effects of one or more known compounds on a cell population in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. Fig. 7K illustrates a step 340 of populating a database
30 with descriptors for known compounds. Such descriptors can be determined from imaging the cell population. However, in some embodiments, descriptors can be derived by measurements and combinations of measurements and the like. Then, in a step 342, descriptors for the unknown compound are determined from imaging a

second cell population. The second cell population has been treated with the unknown compound. Then, in a step 344, a relationship between the descriptors determined from the unknown compound with the descriptors determined from the known compounds can be determined. Finally, in a step 346, an inference can be made about the unknown compound based upon the descriptors of the known compounds from the relationship determined in step 344.

Accordingly, the present invention provides a novel database design. In a particular embodiment according to the present invention, a method for providing a database comprises measurement of a potentially large number of features of one or more sub-cellular morphometric markers. Markers can be from any of a large variety of normal and transformed cell lines from sources such as for example, human beings, fungi, or other species. The markers can be chosen to cover many areas of cell biology, such as, for example markers comprising the cytoskeleton of a cell. The cytoskeleton is one of a plurality of components that determine a cell's architecture, or "cytoarchitecture". A cytoarchitecture comprises structures that can mediate most cellular processes, such as cell growth and division, for example. Because the cytoskeleton is a dynamic structure, it provides a constant indication of the processes occurring within the cell. The cytoarchitecture of a cell can be quantified to produce a one or more scalar values corresponding to many possible cellular markers, such as cytoskeleton, organelles, signaling molecules, adhesion molecules and the like. Such quantification can be performed in the presence and absence of drugs, peptides, proteins, anti-sense oligonucleotides, antibodies, genetic alterations and the like. Scalar values obtained from such quantification can provide information about the shape and metabolic state of the cell.

In a presently preferred embodiment, scalar values can comprise morphometric, frequency, multi-dimensional parameters and the like, extracted from one or more fluorescence images taken from a number of cellular markers from a population of cells. Two or more such scalar values extracted from a plurality of cell lines and markers grown in the same condition together comprise a unique "fingerprint" or descriptor that can be incorporated into a database. Such cellular descriptors will change in the presence of drugs, peptides, proteins, antisense oligonucleotides, antibodies or genetic alterations. Such changes can be sufficiently unique to permit a correlation to be drawn between similar descriptors. Such

correlations can predict similar properties or characteristics with regard to mechanism of action, toxicity, animal model effectiveness, clinical trial effectiveness, patient responses and the like. In a presently preferred embodiment, a database can be built from a plurality of such descriptors from different cell lines, cellular markers, and compounds having known mechanisms of action (or structure, or gene response, or toxicity).

The present invention also provides database and descriptor comparisons according to other embodiments. In a particular embodiment according to the present invention, measurement of scalar values or features can provide predictive information. A database can be provided having one or more "cellular fingerprints" comprised of descriptors of cell substance interactions of drugs having known mechanisms of action with cells. Such descriptors can be compared using a plurality of techniques, such as a technique of creating "phylogenetic trees" of a statistical similarity between the descriptors from various drugs. In a present embodiment, scalar, numeric values can be converted into a nucleotide or amino acid letter. Once converted into a corresponding nucleotide representation, the descriptors can be analyzed and compared using software and algorithms known in the art for genetic and peptide sequence comparisons, such as GCG, a product of Genetics Computer Group, with company headquarters in Madison WI. In an alternative embodiment, numeric values for the fingerprints can be used by comparison techniques. A phylogenetic tree can be created that illustrates a statistical significance of the similarity between descriptors for the drugs in the database. Because the drugs used to build the initial database are of known mechanism, it can be determined whether a particular scalar value in a descriptor is statistically predictive. Finally, a compound fingerprint with no known mechanism of action can be queried against the database and be statistically compared and classified among the drugs in the database that the compound most resembles.

In a particular embodiment, relationships between measured morphometric properties and features of images and physiological conditions can be determined. Relationships can include, for example, treatment of different cell lines with chemical compounds, or comparing cells from a patient with control cells, and the like. In a presently preferable embodiment, a clustering can be performed on acquired image descriptors. Some embodiments can comprise statistical and neural

network - based approaches to perform clustering and comparisons of various descriptors. The foregoing is provided as merely an example, and is not intended to limit the scope of the present invention. Other techniques can be included for different types of data. In some embodiments, clustering and comparing can be performed on features extracted from cell images. In a presently preferable embodiment, procedures for comparisons and phylogenetic analysis of biological sequences can be applied to data obtained from imaging cells.

Select embodiments comprising such approaches enable the use of a broad array of sophisticated algorithms to compare, analyze, and cluster gene and protein sequences. Many programs performing this task are known to those of ordinary skill in the art, such as for example, the program Phylip, available at <http://evolution.genetics.washington.edu/phylip.html>, and other packages listed at <http://evolution.genetics.washington.edu/phylip/software.html>. However, select embodiments according to the present invention can comprise a technique of statistical classification, statistical clustering, distance based clustering, linear and non-linear regression analysis, self-organizing networks, and rule-based classification.

Embodiments can perform such analysis based upon factors such as numerical value, statistical properties, relationships with other values, and the like. In a particular embodiment, numbers in a numerical descriptor can be substituted by one or more of nucleic acid or amino acid codes. Resulting "pseudo-sequences" can be subjected to analysis by a sequence comparison and clustering program.

Other types of databases can also be provided according to other embodiments. The database includes details about the properties of a plurality of standard drugs. When the descriptor of a test compound is compared to the database, predictions about the properties of the test compound can be made using any known property of the other compounds in the database. For example, properties about a compound in the database could include structure, mechanism of action, clinical side effects, toxicity, specificity, gene expression, affinity, pharmacokinetics, and the like. The descriptor of a compound of unknown structure from a natural products library could be compared to the descriptors of compounds with known structure and the structure could be deduced from such a comparison. Similarly, such information could lead to better approaches to drug discovery research including target validation

and compound analogizing, as well as pre-clinical animal modeling, clinical trial design, side effects, dose escalation, patient population and the like.

According to the present invention, databases can be integrated with and complementary to existing genomic databases. Differential genomic expression strategies can be used for drug discovery using database technology. In one particular embodiment, cell data and cellular response data can be associated with a genetic expression profile assay to form a single assay. Live cells expressing fluorescence markers can be treated with a drug, imaged and analyzed for morphometry; and then analyzed for mRNA for expression. Such embodiments can provide rapid development of tools to link cellular behavior with functional genomics.

Database methods according to the present invention can be used to predict gene function and to assist in target validation. Databases that include genetic diversity, i.e., having cellular descriptors from cells of differing genetic backgrounds (tumor, tissue specific, and gene knock out cell lines), can provide the capability to compare cells of unknown genetic background to those in the database. Similarly, the descriptor of an unknown cellular portion in the presence of multiple drugs can be queried against the descriptors of the known markers in the database. For example, if an unknown gene is tagged with Green Fluorescent Protein (GFP), the database may be used to identify the cellular portions for which that unknown gene encodes.

According to the present invention, target validation and specialized cell-based assay screening can be performed using database systems and methods to serve as a universal high-throughput cell-based assay that can evaluate the molecular mechanism of drug action. As new genes are isolated and identified, a large collection of available gene-based knowledge is becoming available. From this large collection of new genes, potential protein targets can be identified using the genomic tools of sequence analysis and expression profiling. However, unless a gene mutation is tightly linked to a disease state, further validation of individual targets is a time consuming process, becoming a bottleneck in drug discovery. Furthermore, robotics and miniaturization are making "High Throughput Screening (HTS)" the industry standard, substantially reducing the time and cost of running a target-based biochemical assay. Therefore, it is now possible to routinely screen large libraries and use a resulting "hit" to validate the target. In such approaches, a specialized cell-based assay would be developed to test hits for each target. Since this often involves

the creation of cell lines expressing new markers, this stage may also become a bottleneck that cannot keep pace with HTS. In addition, these cell-based assays may not be amenable to high-throughput screening, making it difficult to test the increasing number of analogs arising from combinatorial chemistry.

5 In a particular embodiment according to the invention, a rapid characterization of large compound libraries for potential use as pharmaceutical products can be provided by predicting properties of compounds that relate to the compounds' potential as bioactive drugs. In many drug discovery situations, virtually millions of compounds can be passed through a HTS assay against a small number of
10 validated targets. These assays produce hundreds to thousands of potential hits. These hits can then be subsequently screened by a pipeline of secondary and tertiary screens to further characterize their specificity, often time completely missing non-specific interactions with other proteins. Techniques according to the present invention can provide a replacement to such screening operations by providing
15 information about cellular accessibility and mechanism of action for the hits coming from a HTS system. Furthermore, it can replace the biochemical HTS assay and allow rapid and accurate identification of attractive compounds from large libraries without an intervening biochemical assay. The cell information can be predictive of whether to continue into an animal model for each compound, and which animal model to
20 pursue.

 The principles of the present specifically contemplate a wide variety of research methodologies, or usage scenarios, implementing these principles. The following discussion of three such scenarios is by way of illustration and not limitation. Study of the principles enumerated herein will render evident to those
25 skilled in the art certain additional methodologies or usage scenarios enabled by the teachings hereof. The present invention specifically contemplates all such modifications. The following description presents some specific embodiments and scenarios that represent a broader use of cellular phenotypic data and characterizations to deduce mechanisms of action and other features of cellular
30 responses to various stimuli. Such procedures generally involve producing a quantitative cellular phenotype based upon two or more cellular attributes and then comparing that phenotype to phenotypes previously stored and indexed. Such

procedures make use of databases or other repositories of biological information. The invention is not limited to the specific embodiments described here.

Considering first the procedure 2000 depicted in Figure 20, a compound has been identified as having a particular cellular activity. See 2004. For example, a compound may be found to inhibit the growth of certain cancer cell *in vitro* by a specific and desired mechanism of action. This may be a particular company's "gold standard."

Next, the compound is analyzed at 2006 in terms of its effect on one or more cell lines. More specifically, the compound is linked, virtually, to a particular phenotype. Two or more values or measures of cellular attributes characterize that phenotype. These attributes are quantified in the context of specific cellular markers.

In one example, the cellular marker is an organelle such as a nucleus or Golgi apparatus. Measured attributes useful for characterizing an associated phenotype include geometric parameters (e.g., size, shape, and/or location of the organelle) and composition (e.g., concentration of particular biomolecules within the organelle).

The phenotype may be characterized by administering the compound of interest to various cell lines and in various concentrations. In each example within this matrix, the attributes of interest are measured. Ultimately, certain phenotypic features (combinations of attribute values) are associated with the compound of interest. These features provide a template for the phenotype.

Next, using the phenotype as identified at 2006, the process identifies other compounds providing similar features. The goal here is to present a list of compounds having a mechanism of action similar to that of the compound that started the process. This allows researchers to identify a mechanism of action, if not already known, for their compound and to draw conclusions based upon their compound's link to other known compounds (which may not be chemically/structurally similar to the compound of interest).

Identifying similar compounds based upon phenotype can take many paths. Most will involve some mathematical basis. For example, the phenotype defined at 2006 can be represented as a fingerprint or vector comprised of multiple scalar values of cellular attributes (as described above). The phenotype representation can then be compared against known phenotypes characterized by the same format

(e.g., they are all characterized as vectors having the same attribute set, but with different values of the attributes). The comparison may be as simple as a Euclidean distance or more sophisticated as a neural network or multivariate statistical correlation.

5 The known compounds and associated phenotypes may be stored as database records or other data structures that can be queried or otherwise accessed as part of the identification procedure. The compounds may also be associated with other relevant data such as clinical toxicity, cellular toxicity, hypersensitivity, mechanism of action, etc. (when available).

10 Compounds found to be sufficiently similar to the starting compound are returned for consideration by researchers. A data processing system may rank such compounds based on degree of similarity to the starting compound. In some cases, the system may even provide similarity scores associated with the listed compounds.

15 Often researchers wish to determine whether their particular compound has clinical or biochemical effects beyond those that they are already aware of. In a typical scenario, the compound of interest was selected based upon its strong binding a target or its stimulation or inhibition of cell growth in a particular cell line. The process associated with 2010 has likely identified the compound of interest as having
20 a particular mechanism of action based on phenotypic similarity to other compounds having a similar mechanism of action. However, within the region of biochemical space, there may be subspaces (characterized by subphenotypes) that correspond to separate properties. For example, within the phenotypic space associated with one mechanism of action, there may be subspaces associated with clinical toxicity,
25 cellular toxicity (likely overlapping the clinical toxicity space), and little or no toxicity. Obviously, a researcher would like to know whether her compound is likely to be toxic.

 Thus, the process 2000 may include characterizing the compound of interest in terms of its distance from (i.e., similarity to) specific phenotypes having
30 known characteristics. In a typical example, the known characteristic is toxicity. This feature allows the researcher to quantify her compound in terms of mechanism of action AND toxicity (or in terms of two or more other relevant properties associated

with phenotype). To allow simple ranking or characterization, compounds of interest may be scored according to a simple or weighted Boolean expression.

A second scenario of interest is depicted in Figure 21. This scenario again defines a phenotype in terms of a quantifiable vector or other measure.

- 5 However, rather than using a compound of interest to generate the phenotype, some other cellular stimulus is used to generate the phenotype.

As shown, a process 2100 begins with receipt of cells of interest. See 2104. In many situations, the cells are produced by a genetic or epigenetic process that affects the expression level or activity of a particular protein. More generally,
10 any cellular stimulus (e.g., radiation level and type, gravity level, magnetic field, acoustic perturbations, etc.) can be used to generate the cell line of interest. Importantly, this stimulus affects the phenotype and can be correlated therewith.

In the context of drug discovery, a gene encoding for a particular target can be genetically knocked out, underexpressed, overexpressed, expressed in a non-
15 native state, etc. This may be accomplished via standard procedures involving genomic modification, translation or transcription apparatus modification (e.g., use of antisense nucleic acids), blocking target activity (using antibodies to a receptor site for example), and the like. These processes will generally affect the phenotype in some quantifiable way. Importantly, they clearly and unambiguously define a cellular
20 phenotype associated with altering the activity of the target protein.

At 2106, the process involves measuring one or more cellular features from the cell line of interest to define/quantify the phenotype. This may be accomplished as described above with reference to 2006. Next, at 2108, the cellular phenotype generated in this manner is used to identify and rank a set of compounds
25 associated with the phenotype. This operation may proceed in the manner of operations 2008 and/or 2010 from Figure 20.

Finally, at 2110, the process clusters the compounds returned at 2108 by a mechanism of action. The operation 2106 has tightly bound a mechanism of action to a phenotype. Various compounds characterized and stored in a system
30 database may be tentatively assigned a mechanism of action or may have no suggested mechanism of action. By matching their virtual phenotype to the phenotype generated at 2106, one can create or strengthen an association between the compounds and mechanism of action relevant to the stimulus at 2104.

Considering now Figure 22, a third scenario is depicted. This scenario again involves using a virtual phenotype to glean information relevant to a mechanism of action or other cellular activity. In this case, assay data from a group of compounds (e.g., a primary or focused library) is used to elucidate a phenotype.

5 As shown, a process 2200 begins by identifying a target protein. See 2204. Then, at 2206, the process involves identifying positive and negative biochemical hits. More generally, this may involve ranking a number of compounds based upon their interaction with the target. In a specific case, the compounds are ranked based upon their binding affinities to or ability to inhibit the enzymatic activity
10 of the target protein.

After the compounds have been characterized in some manner based upon their interaction with the target, they are used to define a cellular phenotype. See 2208. Generally, the techniques to accomplish are the same as described with reference to operation 2006 of Figure 20. In this case however, one may obtain a
15 strong correlation between mechanism of action (involving the target) and phenotype by using multiple of the compounds identified at 2206. For example, some of the "best hits" may be administered to cell lines in various concentrations. And some of the least effective compounds may also be administered. Cellular attributes that are more strongly exhibited with increasing concentration of the best hits (and not
20 exhibited or exhibited only weakly upon administration of the negative hits) can be used to define the virtual phenotype. In a related approach, compounds having widely varying levels interaction with the target are administered to cells. Those cellular attributes that vary linearly or at least monotonically with the degree of interaction between the target and compound represent attributes that can be used to define the
25 virtual phenotype.

After the cellular phenotype has been defined, previously characterized compounds may be clustered with that phenotype. See 2210. As with operation 2110 of Figure 2, this may create or strengthen an association between a mechanism of action and various compounds in a database.

30 Finally, and optionally, procedure 2200 may provide a "higher resolution" mechanism of action for the compounds identified at 2206. See 2212. Presumably interaction with the target suggests a specific mechanism of action or at least some aspect of a mechanism of action. However, a given target may participate

in a larger cellular mechanism of action – unknown to researchers. Further, a compound may that binds with the target may participate in multiple mechanisms of action – some of which do not involve the target. By linking the target (and its positive hits) to a particular phenotype, some of these additional cellular level activities can be elucidated. The defined phenotype may have been previously identified as associated with other mechanisms of action or higher resolution mechanisms of action. Thus, the phenotype identified at 2208 can be leveraged to generate a higher resolution mechanism of action at 2212.

As suggested in the above discussion, compounds and associated phenotypes may be stored as database records. Such databases can take on many flavors. In one example, a database includes various pieces of information relevant to oncology. Such database may include numerous compounds classified by cellular phenotype, mechanism of action, toxicity, etc. More specifically, the database may include data on commercially available compounds clustered by cellular phenotypes corresponding to mechanisms of action. Further the databases of interest may extended or combined (via standard relational tables and algebra for example) to include additional data such as pharmacology data, cellular genomics data, gene expression data, protein expression data, etc. In a specific example, the database includes measurements made on a subset of the NCI60 cell lines, using DNA, Golgi apparatus, and/or microtubules as markers for defining the phenotypes. Other data includes dosage response information, variation in effect over time, etc. The compounds populating the database could include known National Cancer Institute oncology study compounds. In a specific embodiment, the compound set includes some or all of the compounds mentioned in the article “A gene expression database for the molecular pharmacology of cancer,” Nature Genetics, 24, pp. 236-244 (March 2000).

Various biological analyses may be conducted to develop additional information for characterizing compound mechanisms of action, etc. For example, a cell count analysis may be used to develop dose response curves, GI 50 data, etc. The cell cycle may also be analyzed to find out how various stages in the cycle vary in response to particular stimuli. The Golgi apparatus may be analyzed to determine whether it is in a normal state, a dispersed state, a diffused state, etc. As another example, tubulin may be analyzed to determine whether it is normal, de-polymerized,

over-polymerized, bundled, etc. Obviously, combinations of such analyses may be performed. For example, properties of the Golgi apparatus or tubulin may be analyzed over one or more cell cycles.

In some embodiments, techniques according to the present invention
5 can provide tools for the later stages of drug development such as clinical trial design and patient management. The properties of known drugs, such as clinical trial and patient response information, will be used in a similar fashion as the pre-clinical information to provide predictions about the properties of novel compounds. Because the human cell is the locus of drug action, a database containing drug-cell interactions
10 will be able to provide predictive value for this aspect of drug development.

Although the above has generally been described in terms of specific hardware, software, and methods, it is understood that many alternatives can exist. In particular, the present invention is not limited to a particular kind of data about a cell, but can be applied to virtually any cellular data where an understanding about the
15 workings of the cell is desired. Thus, in some embodiments, the techniques of the present invention could provide information about many different types or groups of cells, substances, and genetic processes of all kinds. Of course, one of ordinary skill in the art would recognize other variations, modifications, and alternatives. Some examples according to the present invention are provided below.

20

EXPERIMENTS

To prove the principle and demonstrate the objects of the present invention, experiments have been performed to determine the effects of manipulations on cell structure using imaging and analysis techniques applied to a variety of
25 situations. These experiments were performed by growing multiple cell lines in the presence of multiple compounds, or substances. Cells were fixed and stained with fluorescent antibodies or labels to multiple cellular portions. One or more images of the cells were then obtained using a digital camera. Descriptors were built by quantifying and/or qualifying patterns of one or more feature from each image in the
30 cell lines under study. A database was built from the descriptors. As the database grows, it should be able to predict the mechanism of action of an unknown drug by comparing its effect with the effects of known compounds or to identify data clusters within large libraries of compounds.

In a first experiment, an automated method to count the number of cells and differentiate normal, mitotic, and apoptotic cells was created.

Approximately, 5,000 HeLa cells were plated per well in a 96 well plate and grown for 3.5 days. The cells were fixed with -20° MEOH for 5 minutes, washed with TBS for 15 minutes, and then incubated in 5 mg/ml Hoechst 33342 in TBS for 15 minutes. Then, 72 images were collected with a 40x objective and 75 ms exposure time.

The analysis was performed on objects that met a certain size criteria that was based on 1) measuring the size of objects in the image that were clearly not cells and 2) excluding the first peak of the area histogram (Fig. 8B values 1-4654).

Histograms of the individual object data were generated for each type of feature. Fig. 8A shows the histogram for average intensity, and Fig. 8B shows histogram data for the area of each object. Fig. 8C shows the scatter plot of the average intensity vs. the area of all of the objects. The pattern of the scatter plot showed an interesting pattern: a large cluster of cells in one region of the graph, with a scattering of object points in other regions. Because mitotic structures are identified as particularly bright objects, most likely due to the biological fact that the chromatin is condensed, the original Hoechst images could be used to identify which cells were either undergoing mitosis, or otherwise looked abnormal. Manual inspection of 917 cells resulted in the classification of each object. Fig. 8D shows a graph where each type of cellular classification is delimited. This graph clearly shows that the mitotic nuclei are brighter than the interphase nuclei. Further, the different phases of the cell cycle can be separated using these two features. Figs. 8E-8F show bar graphs of the average and standard deviations of the areas and average intensities for each cell classification type. These graphs show that interphase nuclei are statistically less bright than mitotic nuclei and that telophase nuclei are statistically smaller than other mitotic nuclei.

Each image was thresholded to an intensity level of 20. A standard area value was set at 9500 pixels. Automated information gathering about all of the objects was done and collected into an Excel spreadsheet (for more information see, section on imaging system). The following information was recorded:

IMAGE NAME
OBJECT #

AREA
STANDARD AREA COUNT
PERIMETER
FIBER LENGTH
FIBER BREADTH
SHAPE FACTOR
ELL. FORM FACTOR
INNER RADIUS
OUTER RADIUS
MEAN RADIUS
AVERAGE INTENSITY
TOTAL INTENSITY
OPTICAL DENSITY
RADIAL DISPERSION
TEXTURE DIFFERENCE MOMENT
EFA HARMONIC 2, SEMI-MAJOR AXIS
EFA HARMONIC 2, SEMI-MINOR AXIS
EFA HARMONIC 2, SEMI-MAJOR AXIS
ANGLE
EFA HARMONIC 2, ELLIPSE AREA
EFA HARMONIC 2, AXIAL RATIO
EFA HARMONIC 3, SEMI-MINOR AXIS

The following results were obtained:

- 1,250 objects were counted
- 201 of those objects has standard area counts > 2 (area > 19000 pixels)
- 195 objects had areas < 6000 pixels
- 1529 objects estimated in total
- 1328 object areas are > 6000 pixels
- The data was reduced to 917 objects that were $6000 < \text{area} < 19000$
- For the 917 objects a scatter plot of area vs. average intensity and a histogram of the average intensity were generated.

- 116 objects that had average intensity intensities > 60 were manually looked at to determine their morphology.
 - Of those 116 objects:
 - 6 were dead or indistinguishable
 - 4 were interphase
 - 30 were prophase
 - 32 were metaphase
 - 24 were anaphase
 - 20 were telophase (10 pairs)
- 10
- 12 prophase objects were missed because of gray scale cut off. (8 of those prophase cells had gray scale values > 57 , as did 7 interphase)
 - 1 telophase object was missed because it was too small (< 6000)
 - 1 prophase object was missed because it was too big (> 1900)
- 15
- 16 mitotic objects were missed because they were parts of objects with standard count > 2 .

In sum, out of 917 single objects, the analysis correctly identified 106 out of 130 mitotic objects, or (81% predictive, 91% of identified mitotics). Out of 917 single objects, the analysis incorrectly identified only 10 non-mitotics as mitotics (1% total, 8% of identified mitotics); 14 mitotics as interphase (1.4% total, 1% interphase). An automated classification system that would automatically assign values to each object using these or other measurement features can thus be developed, utilizing the principles set forth herein.

In a second experiment, the effects of Taxol on MDCK cells and the different types of morphological effects were observed. A plurality of MDCK cells grown in 96 well plates were treated with Taxol for 4.5 hours at different concentrations (10 uM-1pM). They were then fixed, labeled with Hoechst, and imaged.

This experiment used a labeling protocol comprising: MEOH fix at – 20°, Wash in PBS, Block in PBS/BSA/Serum/Triton-X 100, Incubate with 5 µg/ml Hoechst 10 minutes, and wash.

Cells were inspected for different morphologies and manually counted at each different drug concentration in one well. Fig. 9 shows example images from each drug concentration and the different types of morphologies and cells are highlighted. Fig. 10 shows the distribution of each morphology within the cell population as a function of drug concentration. The higher the concentration of Taxol, the larger proportion of cells underwent apoptosis, and the fewer number of normal mitotic cells were detected.

In a third experiment, the purpose was to determine whether the automated analysis methods developed in the first experiment can detect differences in Hoechst morphology in the presence of 6 known compounds at one concentration and exposure time in one cell line. In this experiment, HeLa cells were separately treated with 6 compounds with known mechanism of action. The quantitative methods described in the first experiment were applied to the Hoechst images.

Approximately 5,000 HeLa cells per well were plated in a Costar black-walled 96 well tissue culture treated plate and left to recover in the incubator for 24 hours. After this time, 10 ug/mL of cytochalasin D (CD), Taxol, hydroxyurea, vinblastine, nocodazole, and staurosporine was added to different wells at a 1:100 addition in DMSO.

The cells were incubated in the presence of drug for 24 more hours. After 24 hours, the cells were removed and fixed as in the first experiment. Then, 9 images per well were collected of the Hoechst staining using a 10x objective.

The low magnification images taken of Hoechst were run through the automated image analysis method described in the first experiment. Plots of the average intensity and area were made of each compound. Fig. 11 shows the scatter plots of the compounds. The scatter plots of each compound are visually distinct. For example, cells treated with CD are smaller than control, and cells treated with Hydroxyurea are larger and brighter. Furthermore, the number of cells per well was very different (data not shown).

The effects of different compounds can be clearly and automatically distinguished by identifying changes in cellular morphology. This method can also be used to count adherent cells.

The next experiment was to develop clustering algorithms that assign statistically meaningful values to the representative two dimensional data shown in Fig. 10, and even more complicated clustering of all of the multidimensional data that can be extracted across one, and multiple images.

A fourth experiment was performed to obtain high magnification images of two markers in the presence of drugs. In this experiment, HeLa cells were treated with 80 generic compounds with known mechanism of action. The quantitative methods described in the first experiment were applied to the Hoechst images.

Approximately 5,000 HeLa cells per well were plated in a Costar black walled 96 well tissue culture-treated plate and left to recover in the incubator for 24 hours. After this time, 10 ug/mL of each compound from the Killer Plate from Microsource Discovery Systems (Gaylordsville, CT) was added to different wells at a 1:100 addition in DMSO. The cells were incubated in the presence of drug for 24 more hours. After 24 hours, the cells were removed and fixed as in the first experiment. In addition to being labeled with Hoechst 33342 (against chromatin), cells were also labeled with 1 unit of rhodamine-conjugated phalloidin (against actin) for 30 minutes.

The 96 well plate was imaged twice. Once, 9 images per well were collected of the Hoechst staining using a 10x objective. After this, one image per well of both the phalloidin and Hoechst staining was collected using a 40x objective.

The resulting high magnification images were analyzed qualitatively and distinct pattern differences were detected in both the Hoechst and phalloidin images. Fig. 12 shows three example images from the experiment. The top row is the Hoechst staining, and the bottom row is the phalloidin staining from the same well. The columns show the images from wells treated with just DMSO (control), cytochalasin D, and Colchicine. The morphology of each marker is different in the presence of each drug. Interestingly, there is an effect in the morphology of the chromatin in the Hoechst image of cytochalasin D, which directly targets the actin cytoskeleton (and thus there is an expected effect in the phalloidin image). Also, there is an effect on the actin cytoskeleton, compared to control, in the presence of colchicine that directly targets the microtubule network.

The low magnification images were analyzed as described in the first experiment, and different patterns were seen in both the average intensity vs. area plots, and in the number of cells per well (data not shown). Thus, changes in patterns of a marker that is "down-stream" from the direct target of a compound are detectable. Automated image analysis protocols for actin and other markers can be developed similarly, again utilizing the principles set forth herein.

A fifth experiment was performed to test quadruple labeling of 9 different cell lines grown in normal conditions. In this experiment, NCI-H460, A549, MDA-MD-231, MCF-7, SK-OV-3, OVCAR-3, A498, U-2 OS, and HeLa cells were plated. Then, the cells were fixed and stained for portions of the each cell known as DNA, tubulin, actin, and Golgi.

The following table summarizes the procedures for this experiment:

Action	Active Ingredient/Notes	Buffer	Vol/ well	Desired Time	Temp
Remove media	NOTE: gently by pipetting, not aspiration				
Fix	4% Formaldehyde	PBS	100µl	20 min	rt
Wash		TBS	100µl	5 min	rt
Wash		TBS	100µl	5 min	rt

Permeablize	0.1% Triton X-100	TBS	100μl	10 min	rt
Permeablize	0.1% Triton X-100	TBS	100μl	10 min	rt
Block	% BSA % Serum Filter sterilize before use	TBS w/azide	100μl	1hr or o/n	rt or 4°C
Primary Antibody	1:1000 dilution of DM1α	TBS + 1% BSA + 0.1% TX-100	50μl	1hr or o/n	rt or 4°C
Wash		TBS	100μl	5 min	rt
Wash		TBS	100μl	5 min	rt
Wash		TBS	100μl	5 min	rt
Fluorescent Stain	FITC lens culinaris 1:500 Rhodamine-Phalloidin 1:500 CY5 goat anti-mouse 1:100	TBS + 1% BSA + 0.1% TX-100	50μl	1 hr.	rt, dark
Wash		PBS	100μl	5 min	rt, dark
Hoechst	1:1000 dilution of 5mg/ml	TBS	100μl	15 min	rt, dark
Wash		PBS	100μl	5 min	rt, dark
Wash		PBS	100μl	5 min	rt, dark
Wash		PBS	100μl	5 min	rt, dark
Store		PBS	200μl	1 month	4°C

Cells were plated out at different densities for 48 hours. Cells were fixed and labeled by the above method. Cells were imaged using an automated imaging system that collected 9 images from each marker using a 10x objective.

Higher magnification images were collected of a few cells for demonstration purposes.

In this experiment, each cell line demonstrated different morphological patterns as determined by phase. For example, A549 cells are much more compacted than OVCAR-3 cells as determined by phase contrast imaging (data not shown). The different fluorescent markers showed even bigger differences between different cell lines. Figs. 13 and 14 show 4 panels of each marker for A549 (Fig. 13) and OVCAR-3 cells (Fig. 14). The markers are Hoechst (upper left), Phalloidin (upper right), Lens culinaris (lower left), and DM1a antibody (lower right). The following table summarizes the qualitative differences between these images:

MARKER	A549	OVCAR3
Hoechst/DNA	small	large
Phalloidin/actin	fuzzy	crisp - many stress fibers
Lens culinaris/Golgi	compact	Disperse/punctate
DM1alpha/Tubulin	perinuclear	evenly distributed

Higher magnification images were taken of the OVCAR3 cells. Fig. 15 shows the same markers at 20x, and Fig. 16 shows the markers at 40x. While the highest magnification images show the most detail, these images illustrate that very little morphological or feature information is lost in the 10x images.

These data exemplify the differences in morphology seen between different cell types. Thus the automated image analysis software can be customized for each marker in each cell type. Different drugs should effect these morphologies differentially.

An automated quantification method for each marker and cell line can be similarly developed.

A sixth experiment was conducted with a more sophisticated software package and to develop more flexible image recognition algorithms. In this experiment, prototype image features extraction was performed using MatLab programming language with image toolbox and SDC morphology toolboxes. Algorithms are being developed that will automatically identify objects on images and

to measure various morphological and feature parameters of these objects. Many different features for each of the cellular markers were acquired.

An example of a MatLab program called "AnalyseDNA" that takes as an input an unlimited number of images, identifies individual objects in these images based on either their intensities, or based on edge-detection algorithms, and extracts a number of morphological and intensity characteristics of these objects. A copy of this program follows:

Listing of the AnalyseDNA.m program and of some of the
supporting subroutines

```

10 function files_analysed = AnalyseDNA(filemask, outpath,
    nx, ny, filter_range, dext, modifier, sfname)
    % AnalyseDNA performs measurements on files of DNA images
    % V1. EV 2-11-99; 2-15-99; 2-16-99
15 %
    % files_analysed = AnalyseDNA(filemask, outpath, nx, ny,
    filter_range, dext, modifier, sfname)
    %
    % PARAMETERS:
20 %     ALL PARAMETERS ARE OPTIONAL
    %
    %     FILEMASK - mask for file names to be analyzed
    INCLUDING PATH(for example c:\images\*.tif)
    %     DEFAULT '*.tif' (all *.tif files in the current
25 directory).
    %
    %     OUTPATH - path to a directory where all the output
    files will be placed.
    %     DEFAULT - output is saved in the same directory
30 which contains images
    %
    %     NX, NY - number of individual images in montage
    images along X and Y axes (DEFAULT 1)
    %

```

```
%    FILTER_RANGE - 3 col-wide array (or[]). Specifies
%    how data is filtered when summary is calculated
%    this parameter internally is passed to GetDNADData
%    and then to GetSummaryData - see these
5 %    functions for details. For example: [2 2 Inf; 6 100
%    8000] will case all raws of data for which
%    values in column 2 are less than 2 and all raws
%    where values in column 6 are less than 100 or
%    more than 8000 to be excluded from all
10 calculations of a summary.
%    DEFAULT - [] (means do not filter, summarize all
%    data)
%
%    DEXT - string. Extension for data files being saved.
15 %    DEFAULT 'dat';
%
%    MODIFIER - this modifier is 'SUMMARY', summary file
%    is created;
%    'SUMMARY ONLY' - only summary is generated,
20 data for individual files are not saved
%
%    sfname - string. File name of a summary file
%    DEFAULT 'summary[date].dat'
%
25 % OUTPUT:
%
%    AnalyseDNA works on image files or montages. For
%    each image file it creates a tab-delimits file of
%    measured
30 %    parameters of all the objects in the montage with
%    the same base name as a montage file and extension
%    specified
```

```
%      by dext parameter (or .dat by default) and file
'errors[date].err' - with the list of files that matched
the
%      filemask but could not be processed.
5 %      If 'summary' or 'summary only' modifier is
specified, it also creates a single file
'summary[date].dat' (or
%      different extension, if specified by DEXT) which
contains summary information for all analyzed files.
10 %
%      ALL OUTPUT FILES are saved in a directory specified
by OUTPATH parameter
%
%      RETURNS *files_analysed* - number of files that have
15 been successfully processed.
%
%      Column designations in the output files are
described in GetDNADData
%
20 % FILE NAME CONVENTIONS
%      AnalysedDNA attempts to identify a number for each
file to identify the file in summary output.
%      It does that by looking for the first space or
underscore, followed by a number and then takes
25 %      as many successive numbers as it can find. If it
fails to identify a number it assigns a
%      default which is -1
%
%
30 % SEE ALSO GetDNADData, GetSummaryData
%
% TO DO      improve error handling in opening and writing
files (GLOBAL error_file ?)
```

```
%      include procedures for writing text headers
into the output files

if nargin > 8
5    error ('Wrong number of input parameters');
end
if nargin > 1
    error ('Wrong number of output parameters: only one
allowed');
10 end

% set defaults
need_summary = 0;
summary_only = 0;
15 use_default_outpath = 0;
datestring = datestr(floor(now));
if nargin == 7      % set default summary file name
    sfname = ['summary' deblank(datestring)]; % extension
will be appended later based on dext
20    if deblank(upper(modifier)) == 'SUMMARY'
        need_summary = 1;
    elseif deblank(upper(modifier)) == 'SUMMARY ONLY'
        need_summary = 1;
        summary_only = 1;
25    else
        error (['Wrong parameter: unknown modifier '
modifier]);
    end
end
30
if nargin == 5
    % default data file extension
    set_dext = 'dat';
end
```

```
    if nargin == 4
        % default filter range
        filter_range = [];
    end
5   if nargin == 3
        ny = 1; % default number of images in montage along Y
    end
    if nargin == 2
        nx = 1;
10  end
    if nargin == 1
        use_default_outpath = 1;
    end
    if nargin == 0
15     filemask = '*.tif'
    end

    % check parameters
    if ( ~ischar(filemask) | ~ischar(dext) | ~ischar(sfname)
20  )
        error('Wrong parameter type: filename, filepath,
dext and sfname should be strings');
    end
    if ( ( size(nx) ~= [1 1] ) | ( size(ny) ~= [1 1] ) )
25     error ('Wrong parameter type: nx and ny should be
scalars (1x1 arrays)');
    end
    if (~isempty(filter_range) & size(filter_range, 2) ~= 3)
        error ('Wrong parameter type: filter range should be
30  [] or 3 - cols-wide array');
    end
    % end testing parameters

    % Generate list of files to process
```

```
datapath = getpath(filemask);
if use_default_outpath == 1
    outpath = datapath;
5  end
if exist(outpath, 'dir') ~= 7
    error(['Path ' outpath, 'not found. Exiting..']);
elseif exist(datapath, 'dir') ~= 7
    error(['Path ' datapath, 'not found. Exiting..']);
10 end

sfname = makefullname(outpath, sfname, dext);
if need_summary == 1
    if exist(sfname, 'file')
15     disp(['File ', sfname, 'already exists!']);
        input ('Press ^C to abort, Enter to delete and
continue');
        delete(sfname);
    end
20 end

flist = FileList(getfname(filemask), datapath);
numfiles = size(flist, 1); % total number of files to
25 process
disp(['About to process ', num2str(numfiles), ' files']);
%DEBUG - commented out "input" to run from Wrod
input('Press ^C to abort, Enter to continue');

30 % main loop where the job gets done:
error_file = makefullname(outpath, ['error' datestring
'.err']);
num_processed = 0;
num_error = 0;
```

```
for i = 1:numfiles
    % first generate file name for a data output file
    current_fullname = flist(i, :); % full name with path
    and extension
5    current_datafile = makefullname(outpath,
    makefname(getbasefname(current_fullname), dext) );

    %extract number from a filename
    fnumber = getfilenumber(current_fullname);
10

    % load an imagefile, record errors
    read_error = 0;
    try
        I = imread(current_fullname);
15        %DEBUG
        disp(['Image file #', num2str(fnumber), '
loaded']);
    catch
        % record file-opening error in an error_file
20        read_error = 1;
        num_error = num_error +1;
        msg = [current_fullname ': ' lasterr];
        add_error_msg(error_file, msg);
    end

25
    % extract and write data to a file in outpath
    if read_error ~=1
        if (need_summary == 0)
            %DEBUG
30            disp(['Starting analysis of file #',
num2str(fnumber), '.']);
            current_data = GetDNADData(I, nx, ny, fnumber);
            %DEBUG
```

```

        disp (['Finished analysis of file #',
num2str(fnumber), '.']);
        %load current_data.mat 'current_data';
        write_data(current_data, current_datafile);
5      else      %summary needed
        %DEBUG
        [current_data, current_summary] = GetDNADData(I,
nx, ny, fnumber, filter_range);
        %load current_data.mat 'current_data';
10      %load current_summary.mat 'current_summary';
        write_summary (current_summary, sfname);
        if summary_only ~= 1
            write_data(current_data, current_datafile);
        end
15      end
    end
end % of the main for loop
num_processed = numfiles - num_error;

20  %=====end function AnalyseDNA()
=====

%=====
=====

25  function result = add_error_msg(filename, msg)
    % adds string MSG to an errorfile FILENAME
    % returns 1 if success, 0 if failure

    err_FID = fopen(filename, 'at');
30  if err_FID == -1
        warning(['Can not open error file ' filename]);
    else
        fprintf(err_FID, '%s\n', msg);
        fclose(err_FID);

```



```

end

%=====end function add_error_masg()
=====

5  %=====
=====

function N = getfilenumber(fname)
% returns the first number extracted from a file name
% (string) or -1 if fails to extract any number
10 numbers = NumbersFromString( getfname(fname) ); % vector
    of all numbers encoded in the name

                                % (but not in the path, even if
                                present)
15 if isempty(numbers)
    N = (-1);    % return -1 if no numbers found in the
    name
    else
        N = numbers(1);
20 end

%===== end function getfilenumber()
=====

25 %=====
=====

function result = write_data(data_array, file_name)
% writes data in a data_array in a tab-delimited ascii
file.
30 % result is 0 if success and -1 if failure
% if file_name exists, overwrites it
result = -1;
try
    fid = fopen(file_name, 'wt');

```

```

        if fid ~= -1
            for k = 1:size(data_array, 1)
                fprintf(fid, '%g\t', data_array(k, :));
                fprintf (fid, '\n');
5         end
        test = fclose(fid);
        result = -1;
    catch
        result = -1;
10    end

%===== end function write_data()
=====

15 %=====
=====
function result = write_summary (s_vector, file_name)
% appends summary vector s_vector to a file_name (ASCII
tab-delimited file).
20 % if file_name does not exist, creates it.
% result is 0 if success and -1 if failure
%
result = -1;
try
25     % debug
        fid = fopen(file_name, 'at');
        result = fprintf(fid, '%g\t', s_vector);
        result = fprintf(fid, '\n');
        result = fclose(fid);
30     result = 0;
    catch
        result = -1;
    end
end

```

```

% ===== end function write_summary()
=====

function Data = GetObjectsData(I, Ilabel)
5 % GetObjectsData returns array measurements of objects in
  image "I" masked by "Ilabel"
  % EV 2-3-99; 2-10-99
  % OData = GetObjectsData(I, Ilabel) returns an array of
  morphological and intensity measurements
10 %   taken from a grayscale image "I". Objects are
  identified on a mask image Ilabel, usually
  %   created by bwlablel()
  % OUTPUT:
  % Each row in the output array OData represents
15 individual object
  % columns contain the following measurements:
  %
  %   1 - Index ("number" of an object);      8 -
  Solidity;
20 %   2 - X coordinate of the center of mass; 9 - Extent;
  %   3 - Y coordinate      "-"; 10 - Total
  Intensity;
  %   4 - Total Area (in pixels);      11 - Avg.
  Intensity;
25 %   5 - Ratio of MajorAxis/MinorAxis;      12 - Median
  Intensity;
  %   6 - Eccentricity;      13 - Intensity of
  20% bright pixel
  %   7 - EquivDiameter;      14 - Intensity of
30 80% bright pixel
  %
  % For details on morphological parameters see information
  on MatLab imfeature();

```

```
% Intensity parameters are either obvious or are
documented in comments in this file.

% Procedures in this file are documented in notebook file
"MATLAB Measuring Nuclei (1) 1-29-98.doc"

5
if (nargin ~= 2)
    error ('function requires exactly 2 parameters');
end
if (nargout ~= 1)
10    error ('function has 1 output argument (array X by
    14)');
end

% finished checking arguments

15
% first collect morphological parameters in a structure
array:
ImStats = imfeature(Ilabel, 'Area', 'Centroid',
    'MajorAxisLength',...
20    'MinorAxisLength', 'Eccentricity', 'EquivDiameter',
    ...
    'Solidity', 'Extent', 8 );

% now convert it into array (matrix) while collecting
25 intensity data for each object:

%preallocate output array:
numobjects = size(ImStats, 1);
OData = zeros(numobjects, 14);
30 %now convert ImStats into array and add intensity data to
it
for k=1:numobjects
    OData(k, 1) = k;
    OData(k, 2) = ImStats(k).Centroid(1);
```

```

        OData(k, 3) = ImStats(k).Centroid(2);
        OData(k, 4) = ImStats(k).Area;
        OData(k, 5) = (ImStats(k).MajorAxisLength) /
        (ImStats(k).MinorAxisLength);
5         OData(k, 6) = ImStats(k).Eccentricity ;
        OData(k, 7) = ImStats(k).EquivDiameter;
        OData(k, 8) = ImStats(k).Solidity;
        OData(k, 9) = ImStats(k).Extent;

10         % now collect and assign intensity parameters from
        image I

        object_pixels = find( Ilabel == k);
        object_area = size(object_pixels, 1); %same as total
15 number of pixels in the object
        object_intensities = double(I(object_pixels)); %
        need to convert to double to do math
        sorted_intensities = sort(object_intensities); %
        will need to get median, 20% and 80% pixels
20         total_intensity = sum(object_intensities, 1);
        avg_intensity = total_intensity / object_area;
        median_intensity = sorted_intensities( floor(
        object_area/2 ) + 1 );
        pix20 = sorted_intensities( floor(object_area*0.2)+1
25 ) ; %brightest pixel among dimmest 20%
        pix80 = sorted_intensities( floor(object_area*0.8)+1
        ) ;

        OData(k, 10) = total_intensity;
30         OData(k, 11) = avg_intensity;
        OData(k, 12) = median_intensity;
        OData(k, 13) = pix20; %brightest pixel among dimmest
        20%

```

```

        OData(k, 14) = pix80; %dimmiest pixel among brightest
    20%
    end %for

5   %===== end function
    GetObjectsData()=====

function Imask = MaskDNA1(I);
10  % MaskDNA1 - generates binary mask for cell nuclei
    through edge detection
    % EV 1-22-99; 2-6-99; 2-10-99
    % Imask = MaskDNA1(I)
    % PARAMETERS
15  %   I - intensity image (grayscale)
    % OUTPUT
    %   Imask - BW image with objects from I
    %
    % For more details see Notebook Matlab_DNA_masking1_1-22-
20  99.doc
    % Uses SDC Morphology Toolbox V0.7

    if (nargin ~= 1)
        error('Wrong number of input parameters');
25  end
    if (nargout ~= 1)
        error('Wrong number of output parameters: one output
        argument should be provided');
    end
30

    Imask = edge(I, 'canny');
    Imask = mmdil(Imask, mmsecross(1));
    Imask = mmero ( mmc1ohole(Imask,mmsecross(1)));

```

```

Imask = mmedgeoff(Imask, mmsecross(1));
% note that mmedgeoff this command removed FILLED OBJECTS
but not touching OUTLINES.
% these outlines can be removed by filtering:
5  Imask = medfilt2(Imask, [5 5]);

%=====end MaskDNA1 =====

```

Given the list of image files or montages of images as an input, this
 10 program creates an individual file for each image that contains the following
 quantitative measurements for all objects identified in the image:

- | | |
|---|------------------------------------|
| 1 - Index ("number" of an object); | 8 - Solidity; |
| 2 - X coordinate of the center of mass; | 9 - Extent; |
| 15 3 - Y coordinate "-"; | 10 - Total Intensity; |
| 4 - Total Area (in pixels); | 11 - Avg. Intensity; |
| 5 - Ratio of MajorAxis/MinorAxis; | 12 - Median Intensity; |
| 6 - Eccentricity; | 13 - Intensity of 20% bright pixel |
| 7 - EquivDiameter; | 14 - Intensity of 80% bright pixel |

20 A fragment of an output for a single file, containing 9 images of cells
 stained for DNA and acquired with a 10x objective. A montage image that was used
 as a source to generate data in A is presented in Fig. 17.

The same program also summarizes measurements across many files
 and performs statistical analysis of the summary data. It creates a summary file with
 25 the following data:

- | | |
|--|---------------------------------|
| 1 - Image file number; | |
| 2 - Average object Area (in pixels); | 3 - STD (standard deviation) of |
| 2; | |
| 30 4 - Avg. of Ratio of MajorAxis/MinorAxis; | 5 - STD of 4; |
| 6 - Avg. Eccentricity; | 7 - STD of 6; |
| 8 - Avg. EquivDiameter; | 9 - STD of 8; |
| 10 - Avg. of Solidity; | 11 - STD of 10; |

- | | |
|---|----------------|
| 12 - Avg. of Extent; | 13 - STD of 11 |
| 14 - Avg. of objects Total Intensity; | 15 - STD of 14 |
| 16 - Avg. of objects Avg Intensity; | 16 - STD of 15 |
| 18 - Avg. of objects Median intensity; | 19 - STD of 18 |
| 20 - Avg. of objects intensity of 20% bright pixel; | 21 - STD of 19 |
| 22 - Avg. of objects intensity of 80% bright pixel; | 23 - STD of 21 |

An example of summary output obtained by running AnalyseDNA against 10 montage files also is shown in Appendix B.

10 A seventh experiment was conducted in order to use sequence analysis algorithms to analyze features of cell images. In this experiment, HeLa cells were treated for 24 hours with several different compounds, and then fixed, and stained with a fluorescent DNA dye. One image of these cells was acquired for each of the treatments and morphometric parameters and features were measured:

15 Resulting measurements were arranged into a string of numbers and reduced to a pseudo- nucleic acid sequence using following rules: At any given position in the sequence a number was substituted by "t" (a code for thymidine) if its value is among highest 25% of the values at the corresponding position in the data set, "g" if it is between 50% and 25%, "c" if it is between 75% and 50%, and "a" if it
20 belongs to lowest 25% of values. Thus one descriptor or sequence was generated per treatment as illustrated in Fig. 18.

Resulting sequences were clustered using an AlignX module commercial software package Vector NTI (<http://informaxinc.com>), which uses a Neighbor Joining algorithm for sequence clustering.

25 The resulting dendrogram is presented in Fig 18. On the dendrogram the closest "leafs" correspond to the closest pseudo-sequences. Interestingly, compounds with similar mechanisms of action cluster together on the dendrogram. Another example of the generation of pseudo-sequences and clustering is shown in Fig. 19.

30 In some embodiments, techniques according to the present invention can provide tools for the later stages of drug development such as clinical trial design and patient management. The properties of known drugs such as clinical trial and patient response information will be used in a similar fashion as the pre-clinical

information to provide predictions about the properties of novel compounds. Because the human cell is the locus of drug action, a database containing drug-cell interactions can be able to provide predictive information for this aspect of drug development.

Although the above has generally described the present invention

5 according to specific systems, the present invention has a much broader range of applicability. In particular, the present invention is not limited to a particular kind of data about a cell, but can be applied to virtually any cellular data where an understanding about the workings of the cell is desired. Thus, in some embodiments, the techniques of the present invention could provide information about many

10 different types or groups of cells, substances, and genetic processes of all kinds. Of course, one of ordinary skill in the art would recognize other variations, modifications, and alternatives.

Appendix A

EV Table 1.doc

Example of the output of AnalyseDNA.m program
(measurements for a single 3 by 3 montage image)

File#	Subimage	object#	X coord.	Y coord.	Area	Arm ratio	Eccentricity	Equidiam	Solidity	Extent	Intensity	Avg. Intensity	Median Intensity	20% pla.	80% pla.
1	1	1	12.2897	152.655	115	1.17291	0.522614	13.5875	0.923567	0.739796	4603	31.7366	34	23	37
1	1	2	16.332	416.032	125	1.40594	0.382471	12.6157	0.905391	0.78125	4604	36.848	31	30	45
1	1	3	20.1073	72.8039	127	1.09845	0.413785	15.0121	0.917098	0.681026	4609	26.9435	29	27	31
1	1	4	21.4816	407.144	43	1.16215	0.418004	11.0928	0.914894	0.787457	3490	85.816	87	105	105
1	1	5	21.0316	181	96	1.10887	0.445194	11.0558	0.888889	0.671329	4502	46.8958	49	38	54
1	1	6	30.7332	359.534	206	2.33106	0.401709	16.1953	0.927978	0.715278	6380	30.9709	73	24	37
1	1	7	32.6279	167.371	89	1.34984	0.471694	10.4451	0.927023	0.711667	4725	47.4119	50	39	56
1	1	8	32.6279	167.371	146	1.25176	0.401495	13.6143	0.929836	0.718918	5415	37.089	40	29	44
1	1	9	37.144	344.031	43	1.40462	0.419542	7.73539	0.870337	0.632738	6667	141.851	162	113	131
1	1	10	49.1078	170.004	232	1.90491	0.451127	17.187	0.852941	0.70707	9872	47.3793	65	33	51
1	1	11	56.0749	176.534	211	1.85704	0.455955	16.7164	0.924685	0.686735	7040	37.8552	33	25	33
1	1	12	52.7755	41.9932	71	1.37673	0.463701	13.6009	0.907409	0.706731	4745	32.415	34	26	39
1	1	13	52.4644	346.854	371	2.73725	0.493553	16.7555	0.873449	0.706612	9318	51.8121	56	43	68
1	1	14	56.4079	282.232	206	1.73782	0.454944	16.3553	0.923367	0.67375	7137	31.6156	37	28	41
1	1	15	57.0648	277.176	108	1.71883	0.437266	20.0567	0.925254	0.701799	4644	47	45	51	
1	1	16	64.1714	333.181	315	1.11194	0.474541	12.5143	0.92052	0.626576	4564	40.0513	43	37	47
1	1	17	65.1409	402.411	270	1.70167	0.474541	12.5143	0.92052	0.626576	4564	40.0513	43	37	47
1	1	18	71.8449	443.13	185	1.71883	0.474541	12.5143	0.92052	0.626576	4564	40.0513	43	37	47
1	1	19	72.4849	132.533	106	1.43159	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	20	78.7377	206.27	172	1.3157	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	21	81.4786	51.5812	117	1.44313	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	22	81.4786	51.5812	117	1.44313	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	23	88.1792	261.534	373	2.17388	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	24	84.1765	361.976	85	1.20749	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	25	88.1408	176.211	143	1.43573	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	26	91.6529	316.324	170	1.36553	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	27	97.7604	217.795	288	1.97357	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	28	96.5841	330.363	117	1.04935	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	29	96.5841	249.402	119	1.27374	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	30	103.073	31.3279	127	1.46815	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	31	103.47	155.507	131	1.3208	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	32	103.556	51.1271	118	1.30339	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	33	121.273	265.09	324	1.70379	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	34	123.532	170.445	141	1.57045	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	35	128.88	40.3355	137	1.75649	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	36	137.083	178.083	244	1.845	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	37	130.802	411.5	164	1.19726	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	38	132.439	352.545	187	1.37505	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	39	128.615	16.6136	13	1.15971	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	40	136.518	209.039	101	1.12013	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	41	134.655	37.0909	33	1.11149	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	42	140.893	102.008	121	1.47787	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	43	144.864	59.8159	272	1.55319	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	44	141.093	424.435	161	1.2071	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	45	151.46	256.924	224	1.08008	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	46	153.688	178.516	141	1.39135	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	47	140.875	342.356	48	1.71531	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	48	146.828	11.7677	196	1.42126	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	49	149.613	136.261	217	1.74718	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	50	176.014	356.414	222	1.41888	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	51	175.748	192.983	114	1.33384	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	52	177.181	210.834	147	1.24127	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	53	178.367	101.524	147	1.24127	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	54	182.4	392.431	130	1.34539	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	55	186.4	242.719	186	1.37168	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	56	189.188	252.719	213	1.67083	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	57	200.742	91.3418	213	1.61944	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	58	199.198	156.725	91	1.04653	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	59	208.43	185.871	244	1.94053	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	60	208.752	70.9435	230	1.49129	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	61	212.584	346.655	147	1.13679	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	62	220.654	20.8237	194	2.47462	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	63	216.548	234.28	183	1.34183	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	64	216.582	293.953	171	1.44427	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	65	217.331	330.721	172	1.72537	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	66	217.348	437.1	201	1.29372	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	67	222.826	157.749	121	1.32461	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	68	245.071	484.948	435	1.20117	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47
1	1	69	238.844	283.188	147	1.40408	0.467941	12.6414	0.92052	0.626576	4564	40.0513	43	37	47

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1	1	217,509	86,7157	224	1,87991	0,646782	16,886	0,89598	0,532164	8822	39,4063	42	30	69
2	1	218,831	372,146	160	1,74661	0,630819	14,273	0,91956	0,74555	5025	31,063	33	25	38
3	1	219,214	413,076	77	1,51646	0,498531	9,90149	0,91956	0,74555	5025	56,3195	59	46	61
4	1	220,945	473,816	163	1,36853	0,377002	11,4062	0,91575	0,7	4770	26,5571	30	23	34
5	1	222,001	538,848	66	1,03149	0,245208	9,167	0,91667	0,679167	4770	43,2277	71	56	80
6	1	223,601	598,348	251	1,85991	0,860038	17,8769	0,886926	0,597619	10500	43,2277	71	56	80
7	1	225,001	658,348	161	1,45084	0,495632	14,3175	0,914733	0,488036	5176	31,0066	33	25	38
8	1	226,945	718,848	161	1,31512	0,449722	11,8882	0,917355	0,720759	4958	31,7658	45	36	50
9	1	228,001	778,848	111	1,58165	0,774829	12,8169	0,909722	0,662722	9623	32,7658	45	36	50
10	1	229,945	838,848	204	2,05613	0,873763	18,1165	0,918519	0,596691	7051	34,2657	35	28	41
11	1	231,509	898,848	287	1,27833	0,427825	15,116	0,87274	0,450931	10520	36,4531	63	42	58
12	1	233,601	958,848	150	1,07212	0,327329	13,8188	0,920245	0,765306	9202	41,3167	63	42	58
13	1	235,001	1018,848	65	1,47318	0,601767	10,4031	0,923513	0,817708	4387	51,4118	55	41	56
14	1	236,945	1078,848	271	1,51001	0,74855	17,1499	0,931632	0,675359	6580	27,1429	59	46	61
15	1	238,001	1138,848	221	1,75168	0,872092	16,7746	0,862881	0,701587	10251	46,3816	69	51	67
16	1	239,945	1198,848	66	1,71017	0,810394	7,65708	0,862881	0,701587	10251	151,87	159	170	185
17	1	241,509	1258,848	145	1,31164	0,481667	13,3875	0,917722	0,553208	4910	31,0649	35	27	39
18	1	243,601	1318,848	192	2,01531	0,869208	15,4533	0,911776	0,193388	5972	31,1042	32	24	35
19	1	245,001	1378,848	187	1,25774	0,60151	8,59348	0,920635	0,157143	5103	62,3418	70	55	74
20	1	246,945	1438,848	159	1,79289	0,951172	14,2283	0,928235	0,18125	5169	32,0943	33	26	39
21	1	248,001	1498,848	150	1,31558	0,481667	13,8188	0,925926	0,18125	5169	32,0943	33	26	39
22	1	249,945	1558,848	147	1,35167	0,727432	22,054	0,913267	0,598746	16117	42,1911	44	32	45
23	1	251,509	1618,848	213	1,71848	0,89165	14,3175	0,913047	0,164667	4966	30,8447	33	24	35
24	1	253,601	1678,848	207	1,35063	0,49165	14,3175	0,913047	0,164667	4966	27,3056	28	27	37
25	1	255,001	1738,848	181	1,48164	0,789304	15,1788	0,937642	0,728745	4915	39,2857	60	41	56
26	1	256,945	1798,848	126	1,48164	0,789304	15,1788	0,937642	0,728745	4915	39,2857	60	41	56
27	1	258,001	1858,848	126	2,00031	0,84651	13,4198	0,887353	0,555728	4958	33,0533	35	25	38
28	1	259,945	1918,848	150	1,7164	0,565642	13,5875	0,917722	0,74359	5048	34,8138	36	28	42
29	1	261,509	1978,848	145	1,7164	0,565642	13,5875	0,917722	0,74359	5048	62,551	65	47	64
30	1	263,601	2038,848	147	1,7164	0,565642	13,5875	0,917722	0,74359	5048	33,7119	36	27	39
31	1	265,001	2098,848	213	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
32	1	266,945	2158,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
33	1	268,001	2218,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
34	1	269,945	2278,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
35	1	271,509	2338,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
36	1	273,601	2398,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
37	1	275,001	2458,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
38	1	276,945	2518,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
39	1	278,001	2578,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
40	1	279,945	2638,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
41	1	281,509	2698,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
42	1	283,601	2758,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
43	1	285,001	2818,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
44	1	286,945	2878,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
45	1	288,001	2938,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
46	1	289,945	2998,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
47	1	291,509	3058,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
48	1	293,601	3118,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
49	1	295,001	3178,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
50	1	296,945	3238,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
51	1	298,001	3298,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
52	1	299,945	3358,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
53	1	301,509	3418,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
54	1	303,601	3478,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
55	1	305,001	3538,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
56	1	306,945	3598,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
57	1	308,001	3658,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
58	1	309,945	3718,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
59	1	311,509	3778,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
60	1	313,601	3838,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
61	1	315,001	3898,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
62	1	316,945	3958,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
63	1	318,001	4018,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
64	1	319,945	4078,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
65	1	321,509	4138,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
66	1	323,601	4198,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
67	1	325,001	4258,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
68	1	326,945	4318,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
69	1	328,001	4378,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
70	1	329,945	4438,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
71	1	331,509	4498,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
72	1	333,601	4558,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
73	1	335,001	4618,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
74	1	336,945	4678,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
75	1	338,001	4738,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
76	1	339,945	4798,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
77	1	341,509	4858,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
78	1	343,601	4918,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
79	1	345,001	4978,848	207	1,71848	0,89165	14,3175	0,915515	0,759375	8192	24,1708	25	19	29
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1	165	662.718	316.962	291	2.5992	0.923028	19.7487	0.318719	0.516026	9565	32.8854	32	47
1	166	655.384	305.437	319	1.51731	0.732012	12.2092	0.313385	0.739323	4473	37.3982	39	48
1	167	650.357	299.335	334	1.173345	0.816925	14.0028	0.823158	0.712863	8038	57.1408	48	49
1	168	645.628	300.199	201	1.68173	0.731868	15.9975	0.836888	0.255539	4815	32.9027	24	50
1	169	662.472	163.039	127	1.2763	0.921371	12.7162	0.916784	0.476075	4819	37.2911	35	51
1	170	665.094	234.437	405	1.23102	0.653467	22.7082	0.894061	0.451791	7681	116.8726	118	52
1	150	655	185	23	1.28727	0.537158	5.4152	0.859152	0.537158	4819	35.8286	31	53
1	151	659	228	1.65352	0.785395	17.0083	0.919355	0.919355	0.690801	8144	55.8286	40	54
1	152	660.291	365.573	117	1.37248	0.885151	12.0083	0.919355	0.919355	4819	38.9145	40	55
1	153	665.261	184.174	21	1.37248	0.885151	12.0083	0.919355	0.919355	4819	38.9145	40	56
1	154	672.05	265.007	199	1.37248	0.885151	12.0083	0.919355	0.919355	4819	38.9145	40	57
1	155	680.109	124.184	138	1.31807	0.666436	15.9127	0.916119	0.731077	7636	38.3818	31	58
1	156	685.215	212.969	181	1.31807	0.666436	15.9127	0.916119	0.731077	7636	38.3818	31	59
1	157	693.315	160.964	159	1.44573	0.730898	12.8159	0.916119	0.731077	7636	38.3818	31	60
1	158	698.139	255.492	197	1.18453	0.516037	15.8716	0.916119	0.731077	7636	38.3818	31	61
1	159	698.107	478.713	122	1.36764	0.849465	12.4514	0.916119	0.731077	7636	38.3818	31	62
1	160	501.194	460.519	107	1.30083	0.919359	11.472	0.916119	0.731077	7636	38.3818	31	63
1	161	16.0133	169.201	150	1.72511	0.916119	11.472	0.916119	0.731077	7636	38.3818	31	64
1	1	13.3492	92.819	126	1.72511	0.916119	11.472	0.916119	0.731077	7636	38.3818	31	65
2	1	12.7436	294.949	78	1.24958	0.386585	3.96557	0.904327	0.731077	7636	38.3818	31	66
2	2	12.7436	294.949	78	1.24958	0.386585	3.96557	0.904327	0.731077	7636	38.3818	31	67
2	3	12.7436	294.949	78	1.24958	0.386585	3.96557	0.904327	0.731077	7636	38.3818	31	68
2	4	12.7436	294.949	78	1.24958	0.386585	3.96557	0.904327	0.731077	7636	38.3818	31	69
2	5	12.7436	294.949	78	1.24958	0.386585	3.96557	0.904327	0.731077	7636	38.3818	31	70
2	6	30.8017	150.013	151	1.5317	0.751283	12.8159	0.916119	0.731077	7636	38.3818	31	71
2	7	32.4316	117.441	151	1.01445	0.62005	15.925	0.916119	0.731077	7636	38.3818	31	72
2	8	32.4316	483.965	23	1.17959	0.916119	11.472	0.916119	0.731077	7636	38.3818	31	73
2	9	46.0945	205.184	159	1.33079	0.715132	12.7185	0.916119	0.731077	7636	38.3818	31	74

EV Table 1.doc

EV Table 1.doc	60	221.098	316.41	123	1.19435	0.546784	12.5143	0.871812	0.737243	45733	36.4537	38	29	44
1	1	231.901	273.699	131	1.47119	0.845252	14.7555	0.927034	0.71725	5687	33.1533	35	26	44
2	2	230.372	280.372	132	1.34435	0.791928	15.0545	0.927034	0.698039	4786	26.4876	28	31	44
3	3	233.316	271.425	133	1.94261	0.860461	19.4748	0.845003	0.870833	9681	37.4866	34	26	44
4	4	243.415	495.715	134	1.71632	0.827387	24.7215	0.833852	0.898655	23364	49.4833	52	40	58
5	5	238.415	495.715	135	1.53935	0.784556	14.2283	0.809571	0.684063	6050	38.0203	39	29	47
6	6	238.415	495.715	136	1.30137	0.784556	11.4372	0.809571	0.744252	5042	43.1215	48	39	56
7	7	248.415	116.485	137	1.75031	0.827491	16.5985	0.913127	0.408332	10444	41.9128	50	39	56
8	8	248.415	107.282	138	1.1782	0.528789	14.4057	0.913127	0.777619	4802	27.4801	31	22	35
9	9	251.401	495.351	139	1.36285	0.678453	13.6809	0.914538	0.765625	5259	35.7755	37	29	64
10	10	252.144	448.273	140	1.78272	0.825943	13.3014	0.914538	0.737222	4692	49.3827	51	39	64
11	11	265.218	448.273	141	1.47315	0.733733	14.2733	0.917855	0.765716	5318	33.2275	35	28	36
12	12	270.308	448.273	142	1.82916	0.846443	25.6318	0.765695	0.855789	21246	44.9884	48	34	60
13	13	273.005	102.587	143	1.82916	0.915131	15.9975	0.927018	0.647231	5194	25.0088	27	19	32
14	14	275.753	337.812	144	1.82916	0.938935	14.7122	0.918018	0.708333	4889	29.7588	30	21	37
15	15	282.75	21.4535	145	1.45318	0.722464	14.3166	0.910127	0.736909	8927	54.7325	59	46	48
16	16	282.75	21.4535	146	1.19703	0.594623	15.0545	0.927228	0.741899	8929	55.2191	56	41	68
17	17	304.731	201.003	147	2.0845	0.879415	15.2476	0.911333	0.927552	4598	21.8132	26	19	30
18	18	305.481	404.117	148	1.24994	0.599945	14.1835	0.914511	0.759615	5031	21.8132	26	19	30
19	19	307.247	325.299	149	1.32279	0.831082	12.7162	0.900709	0.641658	4575	25.0716	37	16	44
20	20	310.581	247.256	150	1.52517	0.755036	17.0555	0.931273	0.651832	10311	27.9802	29	21	34
21	21	316.636	202.312	151	1.82819	0.731958	14.7123	0.927397	0.726804	10311	45.0242	41	34	55
22	22	320.641	215.509	152	1.37645	0.681862	17.0382	0.938272	0.76	10187	28.4136	24	23	55
23	23	320.641	215.509	153	1.36105	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
24	24	320.641	215.509	154	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
25	25	320.641	215.509	155	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
26	26	320.641	215.509	156	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
27	27	320.641	215.509	157	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
28	28	320.641	215.509	158	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
29	29	320.641	215.509	159	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
30	30	320.641	215.509	160	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
31	31	320.641	215.509	161	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
32	32	320.641	215.509	162	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
33	33	320.641	215.509	163	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
34	34	320.641	215.509	164	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
35	35	320.641	215.509	165	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
36	36	320.641	215.509	166	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
37	37	320.641	215.509	167	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
38	38	320.641	215.509	168	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
39	39	320.641	215.509	169	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
40	40	320.641	215.509	170	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
41	41	320.641	215.509	171	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
42	42	320.641	215.509	172	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
43	43	320.641	215.509	173	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
44	44	320.641	215.509	174	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
45	45	320.641	215.509	175	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
46	46	320.641	215.509	176	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
47	47	320.641	215.509	177	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
48	48	320.641	215.509	178	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
49	49	320.641	215.509	179	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
50	50	320.641	215.509	180	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
51	51	320.641	215.509	181	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
52	52	320.641	215.509	182	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
53	53	320.641	215.509	183	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
54	54	320.641	215.509	184	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
55	55	320.641	215.509	185	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
56	56	320.641	215.509	186	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
57	57	320.641	215.509	187	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
58	58	320.641	215.509	188	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
59	59	320.641	215.509	189	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
60	60	320.641	215.509	190	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
61	61	320.641	215.509	191	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
62	62	320.641	215.509	192	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
63	63	320.641	215.509	193	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
64	64	320.641	215.509	194	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
65	65	320.641	215.509	195	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
66	66	320.641	215.509	196	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
67	67	320.641	215.509	197	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
68	68	320.641	215.509	198	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
69	69	320.641	215.509	199	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
70	70	320.641	215.509	200	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
71	71	320.641	215.509	201	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
72	72	320.641	215.509	202	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
73	73	320.641	215.509	203	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
74	74	320.641	215.509	204	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
75	75	320.641	215.509	205	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
76	76	320.641	215.509	206	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
77	77	320.641	215.509	207	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
78	78	320.641	215.509	208	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51
79	79	320.641	215.509	209	1.37645	0.954714	16.9633	0.846625	0.934615	4787	46.4788	46	37	51

EV Table I.doc

1	1	136	483.48	366.619	329	2.80072	0.931055	19.5315	0.89521	0.48891	1.065	47.0001	50	38	57
2	2	137	488.732	366.763	190	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
3	3	138	492.310	371.267	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
4	4	139	496.310	376.018	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
5	5	140	500.310	380.769	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
6	6	141	504.310	385.520	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
7	7	142	508.310	390.271	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
8	8	143	512.310	395.022	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
9	9	144	516.310	399.773	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
10	10	145	520.310	404.524	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
11	11	146	524.310	409.275	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
12	12	147	528.310	414.026	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
13	13	148	532.310	418.777	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
14	14	149	536.310	423.528	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
15	15	150	540.310	428.279	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
16	16	151	544.310	433.030	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
17	17	152	548.310	437.781	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
18	18	153	552.310	442.532	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
19	19	154	556.310	447.283	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
20	20	155	560.310	452.034	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
21	21	156	564.310	456.785	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
22	22	157	568.310	461.536	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
23	23	158	572.310	466.287	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
24	24	159	576.310	471.038	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
25	25	160	580.310	475.789	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
26	26	161	584.310	480.540	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
27	27	162	588.310	485.291	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
28	28	163	592.310	490.042	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
29	29	164	596.310	494.793	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
30	30	165	600.310	499.544	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
31	31	166	604.310	504.295	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
32	32	167	608.310	509.046	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
33	33	168	612.310	513.797	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
34	34	169	616.310	518.548	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
35	35	170	620.310	523.299	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
36	36	171	624.310	528.050	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
37	37	172	628.310	532.801	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
38	38	173	632.310	537.552	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
39	39	174	636.310	542.303	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
40	40	175	640.310	547.054	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
41	41	176	644.310	551.805	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
42	42	177	648.310	556.556	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
43	43	178	652.310	561.307	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
44	44	179	656.310	566.058	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
45	45	180	660.310	570.809	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
46	46	181	664.310	575.560	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
47	47	182	668.310	580.311	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
48	48	183	672.310	585.062	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
49	49	184	676.310	589.813	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
50	50	185	680.310	594.564	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
51	51	186	684.310	599.315	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
52	52	187	688.310	604.066	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
53	53	188	692.310	608.817	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
54	54	189	696.310	613.568	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
55	55	190	700.310	618.319	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
56	56	191	704.310	623.070	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
57	57	192	708.310	627.821	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
58	58	193	712.310	632.572	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
59	59	194	716.310	637.323	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
60	60	195	720.310	642.074	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
61	61	196	724.310	646.825	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
62	62	197	728.310	651.576	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
63	63	198	732.310	656.327	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
64	64	199	736.310	661.078	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
65	65	200	740.310	665.829	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
66	66	201	744.310	670.580	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
67	67	202	748.310	675.331	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
68	68	203	752.310	680.082	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
69	69	204	756.310	684.833	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
70	70	205	760.310	689.584	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
71	71	206	764.310	694.335	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
72	72	207	768.310	699.086	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
73	73	208	772.310	703.837	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
74	74	209	776.310	708.588	195	1.75129	0.921058	15.5316	0.835856	0.666667	9186	46.2421	50	35	57
75	75	210	780.310	713.339	195	1.75129	0.921058	15.5316	0.835856	0.666667	9				

74	270.971	433.427	177	1.27996	0.624112	15.0121	0.917098	0.743697	9749	55.0191	55	42
75	271.484	472.443	225	1.27988	0.815864	16.77146	0.917146	0.813689	9810	43.3172	45	33
76	273.472	545.875	291	1.27975	0.91561	16.2377	0.902997	0.418048	9810	47.1635	48	36
77	281.91	595.144	408	2.07161	0.701761	0.741915	0.510435	151333	31.0102	32	24	
78	282.311	590.874	344	1.86644	0.844395	13.5006	0.9	0.686667	1827	51.3547	56	42
79	288.331	758.004	549	2.45553	0.910894	17.8055	0.778195	0.523109	10486	42.1124	43	33
80	280.055	492.405	199	1.27983	0.589177	15.7526	0.921951	0.75	4743	25.0552	27	28
81	283.658	525.6071	112	1.41137	0.741889	11.9416	0.918889	0.422222	4842	47.2321	47	36
82	286.847	776.171	293	1.27224	0.531235	19.1431	0.951299	0.820728	13463	45.9468	47	36
83	281.768	464.0877	114	1.53691	0.749327	12.5651	0.939394	0.480909	4599	48.1919	51	39
84	294.161	369.429	124	1.51031	0.658521	12.8159	0.902098	0.716667	6258	48.5116	51	39
85	296.434	349.49	129	1.32034	0.534531	17.2609	0.902098	0.716667	8552	36.5317	38	28
86	299	473.054	233	1.26718	0.720331	11.9848	0.946875	0.428079	5057	47.708	47	36
87	299.08	329.593	113	1.26718	0.821398	13.4028	0.946875	0.428079	5057	45	47	35
88	305.782	148.492	214	1.26718	0.720331	11.9848	0.946875	0.428079	5057	45	47	35
89	308.427	376.07	118	1.26718	0.821398	13.4028	0.946875	0.428079	5057	36.6094	28	28
90	310.453	401.0355	322	1.26718	0.821398	13.4028	0.946875	0.428079	5057	30.0155	29	21
91	310.53	458.408	473	1.26718	0.821398	13.4028	0.946875	0.428079	5057	30.0155	29	21
92	317.21	754.7	109	1.67571	0.798493	11.2938	0.908001	0.714786	1875	48.75	51	38
93	310.3	754.7	109	1.67571	0.798493	11.2938	0.908001	0.714786	1875	48.75	51	38
94	316.394	172.411	100	1.70651	0.810332	11.7065	0.919655	0.807409	4791	43.9541	44	34
95	319.01	159.242	95	1.4078	0.718907	11.2272	0.904763	0.692208	4732	47.798	49	39
96	320.597	714.444	134	1.4078	0.718907	11.2272	0.904763	0.692208	4732	47.798	49	39
97	330.347	267.994	159	1.17311	0.520073	13.0614	0.905156	0.412121	4915	36.6192	36	28
98	333.319	327.473	21	2.02721	0.718462	16.5355	0.908155	0.47375	10198	37.431	40	29
99	333.177	395.24	192	1.17139	0.81194	11.7986	0.924731	0.754166	9942	57.8023	60	18
100	333.177	395.24	192	1.17139	0.81194	11.7986	0.924731	0.754166	9942	57.8023	60	18
101	337.364	104.017	172	1.40693	0.501534	13.0331	0.934611	0.795466	5241	39.406	40	31
102	340.402	360.612	134	1.53582	0.701539	12.5551	0.916519	0.487216	5215	42.0565	45	33
103	342.452	360.048	174	1.40779	0.821814	15.0847	0.939473	0.619371	5078	26.3487	29	22
104	351.184	265.709	119	1.78123	0.581648	14.7355	0.947059	0.925611	5106	31.7143	32	24
105	349.87	265.707	161	1.71935	0.581648	14.7355	0.947059	0.925611	5106	31.7143	32	24
106	355.373	192.499	206	1.56443	0.82117	14.1515	0.918623	0.43275	7196	37.8442	38	29
107	362.141	126.473	357	1.5789	0.718464	12.7053	0.919658	0.486576	19182	56.2913	57	40
108	364.342	350.4	170	1.28705	0.629539	13.0711	0.915578	0.715697	4816	57.2919	57	40
109	368.919	285.978	46	1.4471	0.729529	12.0711	0.915578	0.715697	4816	57.2919	57	40
110	380.846	738.11	327	1.16149	0.729529	12.0711	0.915578	0.715697	4816	57.2919	57	40
111	381.021	50.5496	131	1.10786	0.82166	12.7182	0.913658	0.755179	4777	31.4412	40	30
112	384.055	301.782	165	1.54783	0.82166	12.7182	0.913658	0.755179	4777	31.4412	40	30
113	386.369	301.782	165	1.54783	0.82166	12.7182	0.913658	0.755179	4777	31.4412	40	30
114	398.813	384.087	134	1.96993	0.74738	15.7357	0.937251	0.800828	9102	59.4061	62	48
115	402.104	171.906	202	2.07428	0.816513	15.7357	0.937251	0.800828	9102	59.4061	62	48
116	407.82	141.787	239	1.7076	0.816513	15.7357	0.937251	0.800828	9102	59.4061	62	48
117	407.82	141.787	239	1.7076	0.816513	15.7357	0.937251	0.800828	9102	59.4061	62	48
118	407.82	141.787	239	1.7076	0.816513	15.7357	0.937251	0.800828	9102	59.4061	62	48
119	407.82	141.787	239	1.7076	0.816513	15.7357	0.937251	0.800828	9102	59.4061	62	48
120	421.21	439.74	100	1.97133	0.816513	15.7357	0.937251	0.800828	9102	59.4061	62	48
121	421.21	439.74	100	1.97133	0.816513	15.7357	0.937251	0.800828	9102	59.4061	62	48
122	428.429	126.659	132	1.71433	0.826053	9.2618	0.918008	0.479197	4008	58.8209	60	47
123	428.429	126.659	132	1.71433	0.826053	9.2618	0.918008	0.479197	4008	58.8209	60	47
124	440.679	30.7469	193	1.33146	0.846287	12.5611	0.927027	0.781015	4590	58.7302	42	31
125	441.316	359.413	266	1.73534	0.846287	12.5611	0.927027	0.781015	4590	58.7302	42	31
126	439.909	324.937	46	1.23646	0.846287	12.5611	0.927027	0.781015	4590	58.7302	42	31
127	439.909	324.937	46	1.23646	0.846287	12.5611	0.927027	0.781015	4590	58.7302	42	31
128	445.932	148.143	95	1.32669	0.846287	12.5611	0.927027	0.781015	4590	58.7302	42	31
129	445.932	148.143	95	1.32669	0.846287	12.5611	0.927027	0.781015	4590	58.7302	42	31
130	441.653	188.412	316	1.21166	0.932769	6.57952	0.85	0.909162	4511	62.0735	25	19
131	447.062	79.8769	130	1.11577	0.932769	6.57952	0.85	0.909162	4511	62.0735	25	19
132	449.534	47.0451	133	1.06714	0.942898	13.0131	0.904762	0.768571	4883	35.2105	36	22
133	450.026	354.31	114	1.28391	0.942898	13.0131	0.904762	0.768571	4883	35.2105	36	22
134	460.24	101.035	224	1.30305	0.942898	13.0131	0.904762	0.768571	4883	35.2105	36	22
135	451.434	476.434	71	1.27893	0.942898	13.0131	0.904762	0.768571	4883	35.2105	36	22
136	462.599	250.088	148	1.36367	0.942898	13.0131	0.904762	0.768571	4883	35.2105	36	22
137	466.043	56.1739	23	1.21672	0.942898	13.0131	0.904762	0.768571	4883	35.2105	36	22
138	465.3	61.85	20	1.20283	0.942898	13.0131	0.904762	0.768571	4883	35.2105	36	22
139	474.911	710.517	87	1.71083	0.846253	50.5248	0.878288	0.666667	2430	121.5	118	149
140	475.395	164.937	159	1.29125	0.939472	14.1815	0.939472	0.77451	4727	43.1379	66	51
141	475.395	164.937	159	1.29125	0.939472	14.1815	0.939472	0.77451	4727	43.1379	66	51
142	482.725	93.054	204	1.77491	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
143	483.752	93.941	201	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
144	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
145	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
146	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
147	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
148	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
149	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
150	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
151	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
152	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
153	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
154	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
155	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
156	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
157	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
158	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
159	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
160	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
161	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6	9319	55.886	156	188
162	480.199	152.449	170	1.77476	0.921609	16.1165	0.908478	0.6				

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1	150	491.486	246.086	35	1.18026	0.513345	6.47538	0.545916	0.833333	35453	101.284	105	89	117
2	151	497.438	246.368	130	2.01206	0.871891	12.8653	0.902718	0.672255	45533	35.0231	36	26	41
3	152	498.438	246.368	128	1.4105	0.591844	12.7682	0.920863	0.711111	4816	37.9335	39	30	46
4	153	504.438	246.368	134	1.4105	0.715017	7.04673	0.906977	0.8125	3680	94.359	56	75	113
5	154	504.438	246.368	132	1.3541	0.635419	13.8116	0.921212	0.730769	4832	31.7895	33	27	40
6	155	504.438	246.368	130	1.32473	0.62327	15.8376	0.929245	0.72983	4940	48.1726	50	70	108
7	156	504.438	246.368	128	1.32473	0.766433	12.1005	0.92	0.746753	4511	39.2261	40	30	48
8	157	504.438	246.368	126	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
9	158	504.438	246.368	124	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
10	159	504.438	246.368	122	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
11	160	504.438	246.368	120	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
12	161	504.438	246.368	118	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
13	162	504.438	246.368	116	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
14	163	504.438	246.368	114	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
15	164	504.438	246.368	112	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
16	165	504.438	246.368	110	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
17	166	504.438	246.368	108	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
18	167	504.438	246.368	106	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
19	168	504.438	246.368	104	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
20	169	504.438	246.368	102	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
21	170	504.438	246.368	100	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
22	171	504.438	246.368	98	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
23	172	504.438	246.368	96	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
24	173	504.438	246.368	94	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
25	174	504.438	246.368	92	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
26	175	504.438	246.368	90	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
27	176	504.438	246.368	88	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
28	177	504.438	246.368	86	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
29	178	504.438	246.368	84	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
30	179	504.438	246.368	82	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
31	180	504.438	246.368	80	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
32	181	504.438	246.368	78	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
33	182	504.438	246.368	76	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
34	183	504.438	246.368	74	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
35	184	504.438	246.368	72	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
36	185	504.438	246.368	70	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
37	186	504.438	246.368	68	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
38	187	504.438	246.368	66	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
39	188	504.438	246.368	64	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
40	189	504.438	246.368	62	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
41	190	504.438	246.368	60	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
42	191	504.438	246.368	58	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
43	192	504.438	246.368	56	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
44	193	504.438	246.368	54	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
45	194	504.438	246.368	52	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
46	195	504.438	246.368	50	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
47	196	504.438	246.368	48	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
48	197	504.438	246.368	46	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
49	198	504.438	246.368	44	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
50	199	504.438	246.368	42	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
51	200	504.438	246.368	40	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
52	201	504.438	246.368	38	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
53	202	504.438	246.368	36	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
54	203	504.438	246.368	34	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
55	204	504.438	246.368	32	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
56	205	504.438	246.368	30	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
57	206	504.438	246.368	28	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
58	207	504.438	246.368	26	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
59	208	504.438	246.368	24	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
60	209	504.438	246.368	22	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
61	210	504.438	246.368	20	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
62	211	504.438	246.368	18	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
63	212	504.438	246.368	16	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
64	213	504.438	246.368	14	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
65	214	504.438	246.368	12	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
66	215	504.438	246.368	10	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
67	216	504.438	246.368	8	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
68	217	504.438	246.368	6	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
69	218	504.438	246.368	4	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
70	219	504.438	246.368	2	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
71	220	504.438	246.368	0	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51
72	221	504.438	246.368	-2	1.3541	0.800492	12.7073	0.931973	0.805882	4042	44.1022	47	35	51

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EV Table 1.doc

1	1	36	246.003	66.7186	334	1.70555	0.810104	70.4219	0.885942	0.703158	15594	46.988	49	38	35
2	1	37	246.016	115.88	165	1.57535	0.712771	13.3875	0.917732	0.697115	4689	32.2359	33	75	55
3	1	38	245.743	266.532	173	1.4535	0.725714	14.8815	0.905159	0.686508	8292	51.711	56	44	44
4	1	39	256.645	297.404	148	1.76551	0.819627	13.7273	0.907975	0.686863	6158	41.6081	48	37	50
5	1	40	245.892	35.5	186	1.69253	0.800796	15.389	0.925373	0.615833	7714	41.4731	44	34	53
6	1	41	267.611	476.84	288	1.69718	0.800108	19.1492	0.957143	0.626087	12451	43.7326	45	33	51
7	1	42	276.033	256.171	152	1.66024	0.798253	13.9116	0.91018	0.703708	6165	41.875	44	34	49
8	1	43	288.878	681.764	173	1.8018	0.831849	12.5143	0.91791	0.803922	4486	36.4715	38	29	41
9	1	44	295.075	642.753	146	1.6836	0.805009	12.6143	0.918239	0.797276	5724	39.2055	40	30	46
10	1	45	300.464	678.541	127	1.79398	0.810231	12.4614	0.917293	0.797186	4489	36.1951	39	30	43
11	1	46	316.606	113.728	254	1.95525	0.859316	17.9834	0.930665	0.574641	10071	39.4528	42	31	47
12	1	47	316.634	410.358	131	1.27616	0.621246	17.9149	0.89326	0.688167	7878	60.145	63	48	73
13	1	48	334.614	418.425	341	1.46951	0.732318	22.3123	0.907851	0.55849	18341	47.9208	47	36	47
14	1	49	334.396	87.0629	141	1.88512	0.847704	17.4915	0.910828	0.752632	1453	32.5185	34	27	38
15	1	50	346.491	100.553	219	2.21268	0.892351	16.6965	0.943964	0.574641	5193	21.7123	24	19	28
16	1	51	346.328	642.77	174	1.41204	0.766034	14.8815	0.925484	0.725	6167	34.597	37	32	44
17	1	52	356.316	136.177	232	1.87911	0.848052	17.187	0.939271	0.658091	3191	22.375	24	18	27
18	1	53	363.332	178.753	265	1.66527	0.798621	16.3687	0.886288	0.750841	11544	41.5623	45	36	52
19	1	54	368.601	52.3114	228	1.73712	0.817484	17.0382	0.919255	0.623333	9119	41.3991	43	33	50
20	1	55	372.972	19.3934	273	1.41274	0.701349	14.4582	0.938276	0.747168	9210	41.2394	44	32	54
21	1	56	374.753	117.607	150	1.29951	0.638622	13.9116	0.943196	0.78125	4817	32.5133	34	27	38
22	1	57	374.303	311.178	152	1.15034	0.494272	13.9116	0.921212	0.71551	4803	31.625	32	24	38
23	1	58	377.8	376.522	135	1.45456	0.724199	12.1005	0.912488	0.711178	4432	38.5824	40	30	46
24	1	59	380.921	365.128	246	1.81817	0.841284	18.4023	0.896312	0.731889	9883	37.1541	39	29	44
25	1	60	384.396	250.143	220	2.04748	0.872617	17.1322	0.916235	0.525	7281	24.0043	38	29	44
26	1	61	385.292	492.442	202	1.46285	0.84372	14.0233	0.915185	0.765152	8562	42.4109	45	34	51
27	1	62	410.504	429.717	138	1.74883	0.820613	13.7074	0.904497	0.620536	6281	30.8705	32	25	36
28	1	63	411.49	466.41	259	1.41752	0.700532	18.1595	0.845355	0.720833	9624	37.146	38	30	45
29	1	64	422.424	135.104	238	1.81888	0.81505	20.1594	0.81592	0.585714	12189	31.7957	35	25	38
30	1	65	423.177	154.422	333	1.41841	0.784244	21.2001	0.82784	0.617813	12182	41.5099	46	35	54
31	1	66	440.34	90.4832	202	1.77459	0.88118	14.0233	0.897378	0.701189	7021	41.584	41	31	45
32	1	67	454.719	235.696	251	2.79159	0.841832	18.0893	0.855312	0.697231	7229	34.2894	38	27	45
33	1	68	452.566	444.405	136	1.58497	0.778016	13.1531	0.925373	0.604964	13025	52.384	55	40	63
34	1	69	464.412	325.218	284	1.22485	0.657225	13.1531	0.90508	0.604964	13025	52.384	55	40	63
35	1	70	468.918	264.443	324	1.42791	0.789633	20.3377	0.915371	0.686314	18203	37.6667	39	29	44
36	1	71	468.918	264.443	324	1.42791	0.789633	20.3377	0.915371	0.686314	18203	37.6667	39	29	44
37	1	72	471.127	211.108	251	1.74241	0.827837	17.8189	0.925794	0.61074	9386	37.3944	39	29	44
38	1	73	465.167	44.9238	131	1.23508	0.585026	11.8982	0.905836	0.706719	4120	37.1171	39	29	44
39	1	74	472.917	140.042	168	1.49841	0.743189	14.6255	0.908108	0.684274	5493	33.8849	36	26	41
40	1	75	472.917	140.042	168	1.49841	0.743189	14.6255	0.908108	0.684274	5493	33.8849	36	26	41
41	1	76	472.917	140.042	168	1.49841	0.743189	14.6255	0.908108	0.684274	5493	33.8849	36	26	41
42	1	77	472.917	140.042	168	1.49841	0.743189	14.6255	0.908108	0.684274	5493	33.8849	36	26	41
43	1	78	480.34	457.404	146	1.088	0.353743	14.1935	0.923877	0.523301	6223	35.3861	43	31	46
44	1	79	483.027	374.5	228	1.19716	0.743189	14.6255	0.908108	0.684274	5493	33.8849	36	26	41
45	1	80	480.34	457.404	146	1.088	0.353743	14.1935	0.923877	0.523301	6223	35.3861	43	31	46
46	1	81	497.325	457.404	146	1.78779	0.827837	17.8189	0.925794	0.61074	9386	37.3944	39	29	44
47	1	82	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
48	1	83	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
49	1	84	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
50	1	85	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
51	1	86	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
52	1	87	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
53	1	88	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
54	1	89	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
55	1	90	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
56	1	91	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
57	1	92	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
58	1	93	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
59	1	94	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
60	1	95	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
61	1	96	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
62	1	97	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
63	1	98	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
64	1	99	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
65	1	100	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
66	1	101	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
67	1	102	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
68	1	103	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
69	1	104	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
70	1	105	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
71	1	106	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
72	1	107	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
73	1	108	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
74	1	109	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
75	1	110	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40
76	1	111	500.488	467.375	124	1.4493	0.820246	12.5451	0.945551	0.632653	4730	35.7258	37	28	40

EV Table 1.doc	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105																												
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
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1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
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1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
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1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
1	2	3	4	5	6	7	8	9</																																																																																																

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106	355.016	312.36	1.76044	0.873412	12.0478	0.897698	0.878374	4181	36.4754	39	45
107	361.714	169.031	1.40135	0.781048	12.8159	0.908451	0.861538	1538	58.4186	46	46
108	370.472	126.786	1.17746	0.48247	11.9416	0.903276	0.717609	4218	37.6607	40	46
109	379.466	101.258	1.21537	0.568327	11.1132	0.898148	0.734888	4148	42.7216	48	46
110	382.737	601.684	1.71489	0.800559	15.4104	0.912195	0.811111	8630	48.1487	48	51
111	388.96	193.188	1.87054	0.815133	19.6463	0.907198	0.613125	11210	38.9967	39	49
112	387.155	331.273	1.75594	0.814985	14.8843	0.915189	0.725	8751	50.3276	52	39
113	384.868	241.441	1.48761	0.804373	17.1559	0.894733	0.653846	4573	37.4544	35	27
114	401.428	302.432	1.54197	0.731095	17.7216	0.865109	0.613333	19684	41.3458	50	32
115	389.517	232.346	2.04845	0.871845	17.7216	0.923664	0.647529	5111	36.3154	38	29
116	416.693	273.039	1.50902	0.748001	27.1118	0.926374	0.601881	9995	26.0266	27	20
117	424.436	30.7862	1.6314	0.760103	14.2293	0.910878	0.80103	4393	28.8805	39	35
118	424.436	16	1.243734	0.584169	11.7808	0.916324	0.762238	3503	35.8013	33	30
119	431.508	178.295	1.38336	0.680966	12.9441	0.921077	0.703333	10217	31.4545	33	25
120	427.646	371.2	1.68431	0.80468	22.4261	0.951007	0.803137	10217	25.8658	24	20
121	435.448	108.709	1.41734	0.713551	10.7441	0.91	0.777778	3764	41.3626	43	33
122	439.949	245.441	1.74057	0.818488	12.9149	0.891156	0.777778	3764	37.7023	34	25
123	441.145	472.391	1.52612	0.755107	11.8145	0.924137	0.785114	4248	38.4091	40	29
124	447.904	87.8989	1.58428	0.775618	15.0545	0.927278	0.65173	4248	25.2497	27	30
125	430.265	488.54	1.37601	0.686912	11.9948	0.941667	0.601145	4125	38.4106	34	29
126	451.785	140.813	1.23762	0.606412	11.672	0.90478	0.710232	3858	38.54	47	31
127	456.12	159.5	1.2749	0.620264	11.2038	0.925976	0.710232	4147	21.2822	37	24
128	461.271	117.674	1.7004	0.808791	17.0131	0.875	0.633333	4147	36.6537	38	28
129	463.58	188.899	1.7079	0.810637	18.0693	0.911148	0.633333	4080	80	79	53
130	472.742	297.325	1.12941	0.644334	11.3201	0.90951	0.715151	4309	29.3172	30	33
131	477.607	334.876	1.23853	0.589592	13.5875	0.921567	0.715151	4309	34.5259	39	30
132	477.543	342.386	1.10393	0.473593	12.153	0.913386	0.715151	4273	39.1743	60	31
133	476.543	22.1651	1.42045	0.710199	11.7806	0.923778	0.710199	4270	30.3581	33	25
134	485.404	60.2073	2.07652	0.876405	14.4503	0.914701	0.700937	5044	101.349	109	45
135	480.397	166.905	1.41513	0.707564	9.95273	0.951351	0.641	6443	39.3716	41	31
136	480.58	254.707	1.53736	0.766698	14.8943	0.935444	0.742359	3881	62.7803	65	51
137	487.806	313.887	1.60841	0.783231	8.88497	0.894551	0.805155	3881	42.7019	43	32
138	496.029	311.5	2.02048	0.871653	16.2317	0.921134	0.721203	4872	41.9192	41	33
139	498.219	7.92709	1.10594	0.677089	11.0558	0.905814	0.721203	4872	36.5555	38	29
140	498.827	358.104	1.55114	0.555584	10.8917	0.920192	0.684595	3878	38.7988	40	32
141	501.108	502.982	1.00155	0.555584	12.1005	0.931192	0.707163	3960	40.1837	42	32
142	512.177	58.4345	1.42734	0.713646	11.2272	0.893009	0.71	3972	62.0104	41	34
143	512.177	58.4345	1.39204	0.695662	11.7064	0.893009	0.71	3972	31.2741	32	21
144	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
145	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
146	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
147	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
148	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
149	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
150	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
151	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
152	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
153	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
154	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
155	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
156	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
157	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
158	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
159	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
160	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
161	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
162	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
163	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
164	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
165	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
166	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
167	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
168	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
169	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
170	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
171	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
172	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
173	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
174	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
175	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
176	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
177	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
178	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
179	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
180	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
181	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
182	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
183	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
184	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
185	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
186	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
187	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
188	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
189	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
190	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
191	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
192	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
193	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
194	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
195	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
196	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
197	512.177	58.4345	1.18415	0.54657	11.0558	0.923077	0.727273	4189	42.44	44	34
198	512.177	58.434									

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1	1	137.631	416.085	116	1.57861	0.77403	12.7573	0.879134	0.7375	4370	37.0319	39	44
2	2	141.049	439.484	162	1.64889	0.84481	15.6353	0.91666	0.827451	4755	24.7656	25	31
3	3	145.252	472.265	248	1.81175	0.933675	17.6788	0.97204	0.859533	5310	31.9178	32	38
4	4	149.455	512.446	341	1.95441	0.99474	19.2988	0.995394	0.727077	5314	30.5955	32	43
5	5	153.658	56.2611	203	1.95361	0.874085	16.0769	0.941188	0.743138	4932	45.2315	32	55
6	6	157.861	219.534	199	1.68464	0.60653	15.3176	0.935644	0.75	4993	25.9616	34	32
7	7	162.062	31.9482	143	1.68464	0.453632	11.4689	0.923197	0.676635	4664	27.5019	28	34
8	8	166.265	745.497	187	2.01862	0.871425	15.4304	0.921182	0.616051	7571	40.4846	42	31
9	9	170.468	33.748	135	1.54998	0.74404	13.1106	0.912162	0.747045	4292	31.7826	33	39
10	10	174.671	386.169	188	2.46016	0.913681	15.7777	0.91093	0.8775	5235	26.4394	37	32
11	11	178.874	501.01	218	2.04299	0.94461	20.4835	0.935821	0.738462	9038	29.2799	31	37
12	12	183.077	180.595	140	1.71516	0.61925	14.273	0.91951	0.683761	4476	29.7935	29	34
13	13	187.280	312.693	177	2.04497	0.874818	15.0121	0.904592	0.706431	6096	34.4607	36	34
14	14	191.483	41.9482	193	1.73535	0.47432	15.6159	0.851278	0.708559	8458	43.879	46	34
15	15	195.686	145.944	143	1.68217	0.601133	12.4935	0.894371	0.647059	4338	30.2657	31	37
16	16	199.889	636.351	171	1.72607	0.453044	12.9149	0.915714	0.729762	4743	31.1577	34	26
17	17	204.092	43.6351	186	1.48471	0.46135	22.7265	0.861143	0.646647	19315	49.8755	52	39
18	18	208.295	236.059	144	1.61497	0.781721	13.9146	0.91212	0.745098	4471	39.0855	30	32
19	19	212.498	466.48	152	1.61497	0.811731	16.2733	0.92054	0.610703	6950	43.0288	44	33
20	20	216.701	34.761	208	2.31997	0.811731	16.2733	0.92054	0.610703	6950	43.0288	44	33
21	21	220.904	74.0298	247	2.07717	0.74481	13.4609	0.91526	0.6125	4256	35.9524	30	30
22	22	225.107	381.666	226	1.61319	0.511034	14.3633	0.933884	0.715542	8913	35.4361	42	30
23	23	229.310	117.696	121	1.80646	0.832822	17.4122	0.93312	0.610559	4781	35.5207	37	41
24	24	233.513	472.545	156	1.52289	0.734197	14.0935	0.864937	0.614338	4765	27.1397	28	21
25	25	237.716	75.6455	131	1.47258	0.801387	11.8145	0.904931	0.733333	6750	57.4515	61	46
26	26	241.919	212.617	206	1.67215	0.811318	14.1953	0.927367	0.734579	7384	21.3701	27	16
27	27	246.122	311.367	160	2.02402	0.646126	14.273	0.870233	0.734579	7384	21.3701	27	16
28	28	250.325	426.811	133	1.78977	0.825716	17.2053	0.914657	0.646773	5538	55.6032	59	44
29	29	254.528	335.378	127	1.50452	0.730763	12.7182	0.927071	0.703135	9241	31.3566	37	21
30	30	258.731	348.239	180	1.32461	0.536017	15.1188	0.927071	0.703135	9241	31.3566	37	21
31	31	262.934	393.9	170	1.61076	0.833135	17.2697	0.914657	0.646773	5538	55.6032	59	44
32	32	267.137	476.726	117	1.66478	0.739719	12.8655	0.927071	0.703135	9241	31.3566	37	21
33	33	271.340	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
34	34	275.543	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
35	35	279.746	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
36	36	283.949	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
37	37	288.152	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
38	38	292.355	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
39	39	296.558	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
40	40	300.761	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
41	41	304.964	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
42	42	309.167	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
43	43	313.370	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
44	44	317.573	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
45	45	321.776	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
46	46	325.979	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
47	47	330.182	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
48	48	334.385	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
49	49	338.588	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
50	50	342.791	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
51	51	346.994	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
52	52	351.197	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
53	53	355.400	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
54	54	359.603	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
55	55	363.806	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
56	56	368.009	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
57	57	372.212	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
58	58	376.415	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
59	59	380.618	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
60	60	384.821	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
61	61	389.024	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
62	62	393.227	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
63	63	397.430	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
64	64	401.633	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
65	65	405.836	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
66	66	410.039	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
67	67	414.242	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
68	68	418.445	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
69	69	422.648	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
70	70	426.851	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
71	71	431.054	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
72	72	435.257	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
73	73	439.460	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
74	74	443.663	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
75	75	447.866	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
76	76	452.069	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
77	77	456.272	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
78	78	460.475	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
79	79	464.678	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
80	80	468.881	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
81	81	473.084	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
82	82	477.287	330.701	127	1.50066	0.745619	12.7182	0.914657	0.646773	5538	55.6032	59	44
83	83	481.490	330.701	127	1.50066	0.745619							

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1	11.1154	387.547	104	1.42586	0.712637	11.5073	0.504346	0.613325	3955	39.0788	39	31	56
2	9.40103	429.712	68	1.2045	0.518981	9.167	0.88	0.713333	3059	46.3485	48	40	53
3	11.6164	443.818	77	1.17632	0.521446	9.0169	0.827731	0.77	2950	38.8312	40	34	46
4	14.4163	456.568	70	1.03128	0.521744	8.4401	0.846076	0.1	1970	56.7143	38	46	68
5	18.1947	45.1316	114	1.265	0.410744	12.0618	0.912	0.710769	1892	34.1404	35	27	41
6	172.59	172.59	100	1.52012	0.753151	11.2038	0.846076	0.711438	3840	38.1	39	31	47
7	25.1163	489.143	140	1.52652	0.753151	11.2038	0.903226	0.613077	4072	29.0857	30	27	38
8	25.8168	433.905	189	1.95343	0.859145	13.2512	0.903226	0.613077	4072	35.1278	31	28	41
9	25.8467	150.616	116	1.27031	0.814445	12.0198	0.919355	0.710769	4004	57.104	40	44	59
10	21.688	113.541	125	1.44331	0.721049	12.6157	0.905787	0.61414	7138	31.6391	32	25	39
11	21.9073	349.684	133	1.23456	0.54442	11.0131	0.817241	0.710769	4208	36.0619	37	29	41
12	28.4991	68.4407	113	1.33116	0.660045	11.5918	0.911328	0.604811	5557	28.4126	29	23	34
13	30.1857	380.439	196	1.80157	0.831803	15.7973	0.903226	0.613077	4072	35.1278	31	28	41
14	36.7162	211.455	143	1.07849	0.371993	13.4935	0.923261	0.727217	11427	36.2131	39	30	46
15	45.0119	37.619	232	1.38806	0.691066	17.9125	0.846076	0.711438	3840	38.1	39	31	47
16	45.7623	123.623	122	1.10508	0.435606	12.4434	0.917293	0.727217	11427	36.2131	39	30	46
17	48.3836	201.731	134	1.40138	0.718897	17.0619	0.917293	0.727217	11427	36.2131	39	30	46
18	50.711	68.5566	212	1.49916	0.744624	16.4284	0.917293	0.727217	11427	36.2131	39	30	46
19	51.1043	461.739	92	1.44718	0.712855	10.423	0.917293	0.727217	11427	36.2131	39	30	46
20	46.0732	162.192	478	1.03578	0.260563	24.67	0.917293	0.727217	11427	36.2131	39	30	46
21	44.5173	121.408	120	1.21481	0.567782	17.3608	0.917293	0.727217	11427	36.2131	39	30	46
22	48.3801	270.181	221	1.5985	0.780155	16.7144	0.917293	0.727217	11427	36.2131	39	30	46
23	44.7156	335.558	156	1.34108	0.763371	14.0935	0.917293	0.727217	11427	36.2131	39	30	46
24	46.7917	50.0764	144	2.23108	0.893926	13.5408	0.917293	0.727217	11427	36.2131	39	30	46
25	47.2043	473	93	1.31433	0.62074	10.8913	0.917293	0.727217	11427	36.2131	39	30	46
26	72.4838	281.915	117	1.44204	0.720491	12.7053	0.917293	0.727217	11427	36.2131	39	30	46
27	72.382	401.976	189	2.01167	0.887693	15.5126	0.917293	0.727217	11427	36.2131	39	30	46
28	76.4716	134.731	114	1.70364	0.809651	13.0419	0.917293	0.727217	11427	36.2131	39	30	46
29	76.4616	109.149	107	1.23867	0.590137	11.672	0.917293	0.727217	11427	36.2131	39	30	46
30	85.2185	188.102	147	2.31787	0.904641	13.4809	0.917293	0.727217	11427	36.2131	39	30	46
31	84.7927	74.7411	113	1.6561	0.511778	11.9948	0.917293	0.727217	11427	36.2131	39	30	46
32	94.0098	208.4	103	1.49478	0.713068	21.6531	0.917293	0.727217	11427	36.2131	39	30	46
33	90.5586	146.101	109	1.51451	0.717418	11.7806	0.917293	0.727217	11427	36.2131	39	30	46
34	86.5458	443.579	273	1.91649	0.833109	18.6639	0.917293	0.727217	11427	36.2131	39	30	46
35	28.1373	243.333	299	1.42403	0.748175	19.5115	0.917293	0.727217	11427	36.2131	39	30	46
36	28.1373	243.333	299	1.42403	0.748175	19.5115	0.917293	0.727217	11427	36.2131	39	30	46
37	28.1373	243.333	299	1.42403	0.748175	19.5115	0.917293	0.727217	11427	36.2131	39	30	46
38	28.1373	243.333	299	1.42403	0.748175	19.5115	0.917293	0.727217	11427	36.2131	39	30	46
39	104.312	441.98	150	1.67096	0.801155	10.8817	0.917293	0.727217	11427	36.2131	39	30	46
40	112.472	222.761	117	1.52261	0.837336	16.1558	0.917293	0.727217	11427	36.2131	39	30	46
41	111.844	344.631	149	1.95261	0.784117	13.8198	0.917293	0.727217	11427	36.2131	39	30	46
42	120.851	611.358	106	1.39004	0.698828	13.7736	0.917293	0.727217	11427	36.2131	39	30	46
43	137.928	192.143	84	1.15403	0.61377	11.4174	0.917293	0.727217	11427	36.2131	39	30	46
44	137.159	654.345	232	1.45957	0.849596	17.9125	0.917293	0.727217	11427	36.2131	39	30	46
45	131.843	178.974	305	2.06123	0.874693	19.7063	0.917293	0.727217	11427	36.2131	39	30	46
46	141.141	215.84	318	1.50812	0.748493	20.1535	0.917293	0.727217	11427	36.2131	39	30	46
47	137.059	742.789	139	1.33113	0.61311	12.1097	0.917293	0.727217	11427	36.2131	39	30	46
48	153.357	17.4886	207	1.75959	0.82639	16.2345	0.917293	0.727217	11427	36.2131	39	30	46
49	153.278	412.183	115	1.65246	0.794105	12.1005	0.917293	0.727217	11427	36.2131	39	30	46
50	159.683	278.456	134	1.43178	0.741056	13.159	0.917293	0.727217	11427	36.2131	39	30	46
51	159.706	91.2227	243	2.00944	0.812777	17.4619	0.917293	0.727217	11427	36.2131	39	30	46
52	159.53	591.024	89	1.4046	0.740198	10.28	0.917293	0.727217	11427	36.2131	39	30	46
53	164.983	500.34	315	1.69183	0.804611	14.9271	0.917293	0.727217	11427	36.2131	39	30	46
54	170.953	257.131	315	1.52018	0.733181	14.5919	0.917293	0.727217	11427	36.2131	39	30	46
55	177.162	54.2575	147	1.20734	0.540216	10.9581	0.917293	0.727217	11427	36.2131	39	30	46
56	178.5	387.358	95	2.70097	0.918922	16.008	0.917293	0.727217	11427	36.2131	39	30	46
57	182.152	125.777	224	1.22471	0.73919	11.3101	0.917293	0.727217	11427	36.2131	39	30	46
58	179.327	459.119	101	1.22471	0.73919	11.3101	0.917293	0.727217	11427	36.2131	39	30	46
59	183.759	190.781	128	1.22471	0.73919	11.3101	0.917293	0.727217	11427	36.2131	39	30	46
60	185.664	303.145	110	1.64455	0.840033	11.4945	0.917293	0.727217	11427	36.2131	39	30	46
61	197.076	139.323	223	1.62485	0.780527	15.4503	0.917293	0.727217	11427	36.2131	39	30	46
62	194.681	214.072	138	2.31277	0.894251	13.2555	0.917293	0.727217	11427	36.2131	39	30	46
63	199.681	455.06	113	1.71704	0.811462	11.3918	0.917293	0.727217	11427	36.2131	39	30	46
64	203.119	282.208	226	1.93002	0.853202	16.3813	0.917293	0.727217	11427	36.2131	39	30	46
65	203.444	35.0173	117	1.24158	0.609669	12.2053	0.917293	0.727217	11427	36.2131	39	30	46
66	206.73	259.208	178	1.65902	0.840137	15.0515	0.917293	0.727217	11427	36.2131	39	30	46
67	210.406	15.8125	128	1.43902	0.728174	12.7662	0.917293	0.727217	11427	36.2131	39	30	46
68	225.178	214.767	163	1.11672	0.473299	24.2213	0.917293	0.727217	11427	36.2131	39	30	46
69	225.178	214.767	163	1.11672	0.473299	24.2213	0.917293	0.727217	11427	36.2131	39	30	46
70	230.979	235.904	146	1.31912	0.65216	11.4303	0.917293	0.727217	11427	36.2131	39	30	46
71	234.613	495.272	146	2.31331	0.903331	20.9981	0.917293	0.727217	11427	36.2131	39	30	46
72	235.102	132.459	255	1.74184	0.820288	18.0188	0.917293	0.727217	11427	36.2131	39	30	46
73	231.124	391.719	89	1.45186	0.724978	10.6451	0.917293	0.727217	11427	36.2131	39	30	46
74	242.343	156.406	251	2.26019	0.894797	17.8769	0.917293	0.727217	11427	36.2131	39	30	46
75	239.34	189.286	162	1.7457	0.820584	14.3618	0.917293	0.727217	11427	36.2131	39	30	46
76	238.514	24.8551	138	1.77073	0.825268	13.2555	0.917293	0.727217	11427	36.2131	39	30	46

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1	1	241.36	45.3128	211	1.49795	0.805148	16.3907	0.546188	0.753571	8135	38.5545	40	30	47
1	1	241.46	300.241	191	1.40918	0.781469	15.5945	0.827184	0.773279	4521	21.4702	25	18	29
1	1	241.56	625.698	149	1.3716	0.760401	14.7116	0.83125	0.716316	4359	29.255	30	22	34
1	1	241.66	379.26	73	1.26745	0.587415	14.0088	0.809244	0.475516	3040	41.9178	33	35	51
1	1	241.76	390.210	156	2.20056	0.800753	14.0955	0.845459	0.5	4763	42.3631	59	47	80
1	1	241.86	248.378	182	1.84198	0.848401	13.7149	0.831126	0.518195	4000	35.8568	37	26	46
1	1	241.96	255.588	266.39	2.23552	0.903116	13.7149	0.89126	0.518195	4000	35.8568	37	26	46
1	1	242.06	203.027	251	2.15737	0.846018	13.7149	0.89126	0.518195	4000	35.8568	37	26	46
1	1	242.16	441.305	131	2.15737	0.846018	13.7149	0.89126	0.518195	4000	35.8568	37	26	46
1	1	242.26	165.613	163	1.28607	0.691448	13.4931	0.82651	0.714702	4196	28.3427	31	23	35
1	1	242.36	165.613	163	1.28607	0.691448	13.4931	0.82651	0.714702	4196	28.3427	31	23	35
1	1	242.46	457.267	268	1.50594	0.7477	18.4724	0.89556	0.708995	8918	31.2161	35	26	40
1	1	242.56	457.267	150	1.60583	0.7477	18.4724	0.89556	0.708995	8918	31.2161	35	26	40
1	1	242.66	40.0113	177	1.12111	0.676231	15.0121	0.814488	0.684444	4722	29.8133	29	23	36
1	1	242.76	417.439	132	1.32613	0.822163	14.7123	0.814488	0.732321	4722	29.8133	29	23	36
1	1	242.86	214.943	105	1.32613	0.822163	14.7123	0.814488	0.732321	4722	29.8133	29	23	36
1	1	242.96	365.215	136	1.32613	0.822163	14.7123	0.814488	0.732321	4722	29.8133	29	23	36
1	1	243.06	44.2279	244	1.32613	0.822163	14.7123	0.814488	0.732321	4722	29.8133	29	23	36
1	1	243.16	83.3276	116	1.25107	0.607565	12.153	0.895225	0.573785	6893	58.4221	41	27	33
1	1	243.26	109.156	178	1.46636	0.73119	15.0967	0.821641	0.727498	6557	37.1899	38	28	46
1	1	243.36	247.71	272	1.32613	0.822163	14.7123	0.814488	0.732321	4722	29.8133	29	23	36
1	1	243.46	479.556	312	1.58325	0.778488	18.9311	0.891429	0.814018	18606	59.4146	67	47	72
1	1	243.56	156.173	235	1.77625	0.826468	18.4134	0.815714	0.770033	9457	31.9493	33	26	38
1	1	243.66	156.173	235	1.77625	0.826468	18.4134	0.815714	0.770033	9457	31.9493	33	26	38
1	1	243.76	216.832	119	1.65657	0.807592	12.3092	0.893771	0.835661	9358	39.1321	44	33	53
1	1	243.86	380.245	102	1.14388	0.660651	11.3561	0.801513	0.653466	4050	34.0216	35	27	41
1	1	243.96	116.445	110	1.14388	0.660651	11.3561	0.801513	0.653466	4050	34.0216	35	27	41
1	1	244.06	356.747	91	1.50007	0.745185	11.8145	0.810714	0.728571	3971	38.9216	40	30	47
1	1	244.16	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	244.26	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	244.36	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	244.46	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	244.56	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	244.66	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	244.76	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	244.86	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	244.96	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	245.06	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	245.16	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	245.26	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	245.36	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	245.46	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	245.56	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	245.66	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	245.76	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	245.86	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	245.96	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	246.06	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	246.16	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	246.26	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	246.36	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	246.46	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	246.56	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	246.66	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	246.76	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	246.86	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	246.96	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	247.06	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	247.16	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	247.26	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	247.36	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	247.46	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	247.56	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	247.66	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	247.76	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	247.86	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	247.96	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	248.06	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	248.16	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	248.26	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	248.36	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	248.46	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	248.56	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	248.66	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	248.76	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	248.86	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	248.96	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	249.06	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	249.16	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	249.26	216.455	121	1.99171	0.864821	10.7841	0.801639	0.651662	4017	36.5182	38	28	44
1	1	24												

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1	7	21.2188	361.175	64	1.29161	0.61372	9.02703	0.07812	0.72723	3642	56.8063	57	44	37
2	8	24.0231	451.946	129	1.82866	0.81321	12.8158	0.121429	0.66158	3872	30.0155	30	23	37
3	9	35.4439	114.293	140	1.45761	0.72755	13.3532	0.108091	0.729167	4313	30.4071	31	23	38
4	10	38.9272	251.374	206	1.6079	0.79493	14.1953	0.119643	0.471632	5014	41.7573	45	35	53
5	11	38.5562	278.849	178	1.06005	0.731755	15.0543	0.127083	0.791111	9196	51.6579	52	38	61
6	12	39.3667	61.32	150	1.17668	0.527089	13.8158	0.120245	0.765506	4709	31.3933	33	25	38
7	13	38.3745	348.388	85	1.17668	0.527089	13.8158	0.120245	0.765506	3980	46.4235	69	39	57
8	14	45.2733	24.1727	139	1.40448	0.703172	13.10316	0.119643	0.471632	4278	30.7642	32	25	36
9	15	51.1316	103.968	181	1.77185	0.835478	15.4304	0.121188	0.618263	4110	23.9879	24	18	29
10	16	55.8904	67.5276	146	1.41539	0.745475	13.6143	0.1274031	0.695238	4258	28.4344	30	22	36
11	17	55.5476	126.373	176	1.31687	0.650318	12.666	0.1274031	0.695238	4084	32.5238	33	25	40
12	18	65.1618	170.866	374	2.47752	0.914573	20.3108	0.15807	0.482143	18200	50.7	52	39	61
13	19	81.8615	292.118	195	1.45938	0.799015	15.757	0.159111	0.714286	9253	46.7821	48	35	57
14	20	63.3444	214.012	180	1.72726	0.825678	15.1388	0.160573	0.714286	8730	48.5	50	37	59
15	21	77.2722	275.178	45	1.42858	0.716165	15.5694	0.160573	0.714286	9364	74.8	74	56	81
16	22	76.2075	362.952	153	1.44916	0.730838	13.9573	0.160573	0.714286	4772	75.7286	74	56	81
17	23	81.9178	94.1466	192	1.14916	0.692694	12.4732	0.160573	0.714286	4772	75.7286	74	56	81
18	24	96.0609	317.476	115	1.25503	0.621756	15.6353	0.160573	0.714286	4772	75.7286	74	56	81
19	25	98.0948	190.518	197	1.14488	0.684908	15.8776	0.160573	0.714286	4772	75.7286	74	56	81
20	26	96.4796	167.951	103	1.17947	0.658936	11.4518	0.160573	0.714286	4772	75.7286	74	56	81
21	27	102.931	121.97	203	2.8642	0.938412	16.0769	0.160573	0.714286	4772	75.7286	74	56	81
22	28	109.951	38.6539	122	1.77624	0.813318	12.4631	0.160573	0.714286	4772	75.7286	74	56	81
23	29	101.027	99.8591	149	1.68447	0.841728	13.7736	0.160573	0.714286	4772	75.7286	74	56	81
30	30	99.8	243.069	130	1.27609	0.621756	12.8655	0.160573	0.714286	4772	75.7286	74	56	81
31	31	104.122	619.385	109	1.17019	0.519347	11.7806	0.160573	0.714286	4772	75.7286	74	56	81
32	32	109.131	516.167	199	1.51357	0.75064	15.9177	0.160573	0.714286	4772	75.7286	74	56	81
33	33	104.156	111.843	115	1.22263	0.672545	10.7047	0.160573	0.714286	4772	75.7286	74	56	81
34	34	110.172	713.843	315	1.22263	0.672545	10.7047	0.160573	0.714286	4772	75.7286	74	56	81
35	35	110.172	713.843	315	1.22263	0.672545	10.7047	0.160573	0.714286	4772	75.7286	74	56	81
36	36	110.172	713.843	315	1.22263	0.672545	10.7047	0.160573	0.714286	4772	75.7286	74	56	81
37	37	116.382	401.511	91	1.27222	0.57861	11.1704	0.160573	0.714286	4772	75.7286	74	56	81
38	38	116.382	401.511	91	1.27222	0.57861	11.1704	0.160573	0.714286	4772	75.7286	74	56	81
39	39	128.94	111.753	103	1.20836	0.610508	11.2772	0.160573	0.714286	4772	75.7286	74	56	81
40	40	128.94	111.753	103	1.20836	0.610508	11.2772	0.160573	0.714286	4772	75.7286	74	56	81
41	41	138.116	195.913	277	1.75132	0.813046	16.78	0.160573	0.714286	4772	75.7286	74	56	81
42	42	138.116	195.913	277	1.75132	0.813046	16.78	0.160573	0.714286	4772	75.7286	74	56	81
43	43	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
44	44	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
45	45	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
46	46	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
47	47	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
48	48	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
49	49	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
50	50	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
51	51	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
52	52	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
53	53	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
54	54	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
55	55	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
56	56	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
57	57	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
58	58	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
59	59	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
60	60	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
61	61	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
62	62	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
63	63	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
64	64	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
65	65	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
66	66	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
67	67	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
68	68	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
69	69	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
70	70	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
71	71	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
72	72	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
73	73	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
74	74	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
75	75	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
76	76	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
77	77	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
78	78	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
79	79	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
80	80	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
81	81	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
82	82	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
83	83	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
84	84	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772	75.7286	74	56	81
85	85	145.814	211.481	101	1.19519	0.627316	15.4304	0.160573	0.714286	4772</				

EV Table 1.doc

93	293.616	101.011	127	1.5886	0.805767	12.7167	0.913669	0.661639	4625	36.4173	37	29	41
94	284.616	410.671	173	1.51516	0.739964	16.8035	0.928134	0.759772	7670	44.3153	41	35	51
95	281.616	289.917	94	1.53154	0.765811	10.3018	0.907376	0.664816	4211	50.131	51	38	56
96	286.316	80.4481	154	1.13333	0.470616	16.0028	0.916687	0.723333	1524	29.1996	20	33	36
97	292.989	305.911	90	1.08188	0.393714	10.7047	0.908091	0.719002	6136	45.9333	46	34	56
98	300.374	47.9316	107	1.37271	0.485058	11.4727	0.920053	0.765288	4067	36.0093	19	29	46
99	302.5	337.755	106	1.53534	0.758839	11.6174	0.938053	0.841327	1706	44.3196	41	36	54
100	310.561	318.784	148	2.38765	0.908093	13.7273	0.920053	0.731971	8927	40.4327	40	65	106
101	305.159	344.711	44	1.29753	0.437205	7.48882	0.864156	0.6875	3746	85.3565	88	61	106
102	313.103	388.246	224	2.05865	0.874084	16.888	0.912266	0.59891	5745	41.7168	41	37	50
103	312.377	242.253	359	2.13587	0.881628	12.3397	0.910784	0.565941	19817	53.2008	56	42	68
104	315.082	332.148	182	1.68477	0.738934	15.2227	0.928571	0.731642	9741	35.0549	35	41	64
105	332.65	51.4728	164	1.12189	0.435902	14.4052	0.910615	0.619216	14361	26.7733	28	30	32
106	327.5	256.005	158	1.26078	0.624816	14.1835	0.923977	0.752781	4988	31.5696	33	25	33
107	327.743	252.568	248	1.44039	0.793938	17.5172	0.895911	0.618211	9916	41.1552	42	31	51
108	342.317	45.2164	351	1.33818	0.461971	13.8658	0.927032	0.725962	4216	26.1393	29	27	31
109	344.143	335.007	28	1.32688	0.460686	5.97082	0.875	0.666667	8659	202.1017	215	166	218
110	356.063	396.314	173	1.60946	0.763613	14.8415	0.935155	0.759772	4659	27.1418	28	21	31
111	355.063	396.314	145	1.37417	0.693684	13.5875	0.928487	0.761659	4823	58.0897	60	47	69
112	318.17	76.9816	217	2.28618	0.898918	16.4271	0.90795	0.618933	4593	21.7553	21	17	22
113	338.408	69.4078	74	1.22379	0.754432	9.81698	0.894818	0.608009	7205	94.0076	99	76	114
114	348.361	67.4555	11	1.13076	0.478395	3.74241	0.785314	0.68875	210	19.0093	19	14	22
115	350.203	202.574	146	1.81956	0.939147	14.5381	0.922272	0.65873	4586	29.0966	30	22	35
116	371.771	706.053	604	1.51498	0.450488	22.4358	0.930876	0.704294	70716	51.2732	52	37	46
117	384.148	245.458	328	1.61555	0.764718	20.4358	0.946104	0.627933	3544	36.6324	37	28	46
118	395.638	462.118	65	1.52246	0.754003	9.37702	0.92619	0.645933	6337	26.7362	20	22	38
119	383.181	491.699	213	1.72925	0.821648	16.4482	0.924087	0.719593	4988	44.1098	51	39	59
120	391.713	651.316	735	1.60311	0.831703	15.1337	0.924171	0.711286	8007	46.1937	48	35	58
121	404.037	405.346	62	1.74935	0.779393	10.2179	0.911111	0.758258	3945	31.1664	31	26	41
122	416.117	114.047	60	1.73356	0.898125	12.5651	0.905109	0.738094	8216	31.366	39	30	46
123	430.317	385.28	726	1.53531	0.72655	17.8412	0.91995	0.718094	4216	51.9644	52	41	61
124	436.472	217.156	250	1.43531	0.679797	9.57661	0.87809	0.72	2710	51.9644	52	41	61
125	432.477	682.136	128	1.21482	0.483784	20.2395	0.913602	0.419481	13119	34.0293	36	27	42
126	455.773	400.5	130	1.41995	0.709932	12.6855	0.935933	0.433333	4951	32.3324	31	26	40
127	416.273	107.954	120	1.48858	0.760788	13.5116	0.935933	0.433333	4951	32.3324	31	26	40
128	479.157	215.248	321	1.77349	0.61919	12.4122	0.903983	0.715976	3981	25.3524	23	28	33
129	408.488	10.1873	49	1.4708	0.732302	7.88865	0.907407	0.737778	4290	120.224	122	92	143
130	485.018	325.565	168	1.6403	0.792485	14.6255	0.933333	0.737778	4290	25.5157	26	32	32
131	485.018	325.565	168	1.6403	0.792485	14.6255	0.933333	0.737778	4290	25.5157	26	32	32
132	488.055	270.544	160	1.36579	0.68109	14.273	0.946766	0.768434	8081	50.5083	53	41	61
133	489.577	41.2515	165	1.43325	0.731835	14.4183	0.931102	0.705128	4206	25.4509	27	40	40
134	488.927	311.45	191	1.39613	0.697829	15.5945	0.927188	0.702306	9350	49.9559	50	38	40
135	492.91	94.3319	212	1.46811	0.854034	16.4394	0.921718	0.708385	15000	35.3774	38	28	35
136	495.135	339.708	192	1.22112	0.457671	15.6353	0.933039	0.761905	8831	45.9948	48	26	32
137	497.009	369.288	178	2.12712	0.881893	15.0345	0.92228	0.706469	6413	36.0727	18	21	24

EV Table 2.doc

Example of the summary output of AnalyseDNA.m program
(summary for 10 3 by 3 montage images)

1	1187	143.912	79.3918	1.59219	0.388735	0.726461	0.133996	14.0412	3.315	0.805327	0.0350365	0.701218	0.075176	6149.26	1456.35	61.539	18.352	62.9414	18.9393
2	1203	14.314	50.5906	22.6594	0.389942	0.727835	0.134442	14.2434	3.43288	0.906571	0.0311453	0.70177	0.0730289	6786.37	3095.1	62.2416	17.0965	63.0245	17.383
3	1205	169.016	86.8722	1.60511	0.389942	0.727835	0.134442	14.2434	3.43288	0.906571	0.0311453	0.70177	0.0730289	6786.37	3095.1	62.2416	17.0965	63.0245	17.383
4	1205	169.016	86.8722	1.60511	0.389942	0.727835	0.134442	14.2434	3.43288	0.906571	0.0311453	0.70177	0.0730289	6786.37	3095.1	62.2416	17.0965	63.0245	17.383
5	1205	169.016	86.8722	1.60511	0.389942	0.727835	0.134442	14.2434	3.43288	0.906571	0.0311453	0.70177	0.0730289	6786.37	3095.1	62.2416	17.0965	63.0245	17.383
6	1205	169.016	86.8722	1.60511	0.389942	0.727835	0.134442	14.2434	3.43288	0.906571	0.0311453	0.70177	0.0730289	6786.37	3095.1	62.2416	17.0965	63.0245	17.383
7	1205	169.016	86.8722	1.60511	0.389942	0.727835	0.134442	14.2434	3.43288	0.906571	0.0311453	0.70177	0.0730289	6786.37	3095.1	62.2416	17.0965	63.0245	17.383
8	1205	169.016	86.8722	1.60511	0.389942	0.727835	0.134442	14.2434	3.43288	0.906571	0.0311453	0.70177	0.0730289	6786.37	3095.1	62.2416	17.0965	63.0245	17.383
9	1205	169.016	86.8722	1.60511	0.389942	0.727835	0.134442	14.2434	3.43288	0.906571	0.0311453	0.70177	0.0730289	6786.37	3095.1	62.2416	17.0965	63.0245	17.383
10	1205	169.016	86.8722	1.60511	0.389942	0.727835	0.134442	14.2434	3.43288	0.906571	0.0311453	0.70177	0.0730289	6786.37	3095.1	62.2416	17.0965	63.0245	17.383

CLAIMS

What is claimed is:

1. A method of predicting a property of a manipulation of cells based
5 upon a descriptor, said method comprising:
 providing a plurality of cells;
 manipulating said plurality of cells;
 capturing a morphological value from said plurality of cells;
 assigning a degree of presence of said morphological value; and
10 storing said morphological value and said degree of presence;
 wherein said descriptor is derived from a first component of a cell and
a second component of said cell, said capturing said morphometric value from said
plurality of cells comprises determining a relationship between said first component
and said second component.
- 15 2. The method of claim 1 wherein said first component and said second
component are selected from a protein, a protein modification, a nucleic acid, a lipid,
a carbohydrate, a subcellular structure and an organelle.
3. The method of 1 wherein said step of manipulation occurs in a manner
selected from a electrical source, a chemical source, a thermal source, a gravitational
20 source, a nuclear source, a temporal source, and a biological source
4. The method of claim 3 wherein said chemical source is selected from a
pharmacological candidate and a drug screening library.
5. The method of claim 1 wherein said morphological value is selected
from a count, an area, a perimeter, a length, a breadth, a fiber length, a fiber breadth, a
25 shape factor, a elliptical form factor, an inner radius, an outer radius, a mean radius,
an equivalent radius, an equivalent sphere volume, an equivalent prolate volume, an
equivalent oblate volume, an equivalent sphere surface area, an average gray value, a
total gray value, and an optical density.
6. The method of claim 1 wherein said degree of presence is
30 multiple of a quantized value.

7. A computer program product for populating a database with manipulated biological information, said computer program product comprising:
- code for providing a plurality of cells in various stages of the cell cycle, said stages of the cell cycle including at least one selected from interphase, G0 phase, G1 phase, S phase, G2 phase, M phase, prophase, prometaphase, metaphase, anaphase, and telophase;
 - code for manipulating said cells in said various stages of cell cycle development to form a plurality of manipulated cells;
 - code for capturing an image of said plurality of manipulated cells;
 - code for determining a descriptor from said image for said manipulated cells;
 - code for populating a database with said descriptor;
 - wherein said image includes a first component of a cell and a second component of said cell; and
 - a computer readable storage medium for holding the codes.
8. The computer program product of claim 7 wherein said first component and said second component are selected from a protein, a protein modification, a nucleic acid, a lipid, a carbohydrate, a sub-cellular structure and an organelle.
9. The computer program product of claim 7 wherein said image is a digitized representation of said plurality of manipulated cells.
11. The computer program product of claim 9 wherein said digitized representation provides a density value of said plurality of manipulated cells.
11. The computer program product of claim 7 wherein said descriptors comprise numeric or logical values.
12. The computer program product of claim 11 wherein said values further comprises a nucleotide.
13. The computer program product of claim 11 wherein said values further comprises an amino acid letter.
14. A system for capturing images of cells or cell structures, the system comprising:
- a cell holder comprising a plurality of sites in a spatial orientation, each of the sites being capable of holding a plurality of cells to be imaged;

an image capturing device coupled to the cell holder, the image capture device being adapted to capture at least one image in at least one of the plurality of sites;

an illumination apparatus comprising a liquid light guide coupled to the plate for highlighting the plurality of cells in a relatively even spatial manner for image capturing purposes;

an image processing device coupled to the image capturing device, the image capturing device being adapted to convert the image into a digital representation; and

a database storage device comprising a database management element coupled to the image capturing device, the database storage device being adapted to retrieve the digital representation of the image from the image processing device and storing the digital representation.

15. The system of claim 14 further comprising a stage comprising a device for moving the cell holder in a spatial direction to traverse across the cell holder in the spatial orientation.

16. The system of claim 14 wherein the illumination apparatus comprises sub-elements, at least one of the sub-elements being positioned away from the image capturing device to prevent a possibility of vibration from the one sub-elements to be transmitted to the image capturing device.

17. The system of claim 14 wherein the digital representation comprises a plurality of regions and objects.

18. The system of claim 14 further comprising a computing device connected between the database storage device and the image processing device.

19. The system of claim 14 wherein the image capturing device comprises a magnification of at least 1X and greater to capture the image of the site.

20. The system of claim 14 wherein the plurality of sites comprises at least 96 sites.

21. The system of claim 14 wherein the liquid light guide characterized as a flexible member that substantially prevents vibration from the an element of the illumination apparatus to be transferred to the image capturing device.

22. The system of claim 14 wherein the spatial direction can be selected from an x-direction, a y-direction, or a z-direction in a Cartesian coordinate system.

23. The system of claim 14 wherein the each of the sites comprises
5 a volume that is sufficient to prevent a solution therein from evaporating in a substantial manner that may influence the image capturing.

24. A method for identifying a mechanism of action for a first compound, the method comprising the steps of:
receiving the first compound;
10 measuring at least one feature of a cellular phenotype to define a target phenotype;
identifying additional compounds providing a feature similar to the feature identified in the measuring step; and
characterizing the first compound in terms of distance from a specific
15 target phenotype having known characteristics.

25. The method of claim 24 comprising the further step of storing the additional compounds and their associated phenotypes in a database.

26. A method for identifying a mechanism of action for a cellular stimulus, the method comprising the steps of:
20 receiving cells of interest;
measuring at least one feature of the cells to define and quantify a target phenotype;
identifying additional compounds providing a feature similar to the feature identified in the measuring step; and
25 characterizing the first compound in terms of distance from a specific target phenotype having known characteristics.

27. A method for identifying information relevant to at least one of a mechanism of action and cellular activity by utilizing assay data to elucidate a phenotype, the method comprising the steps of:
30 identifying a target protein;
identifying positive and negative biochemical hits related to the target protein;
defining the target phenotype utilizing the positive and negative hits;

identifying other compounds providing similar features; and
characterizing the first compound in terms of distance from a specific
target phenotype having known characteristics.

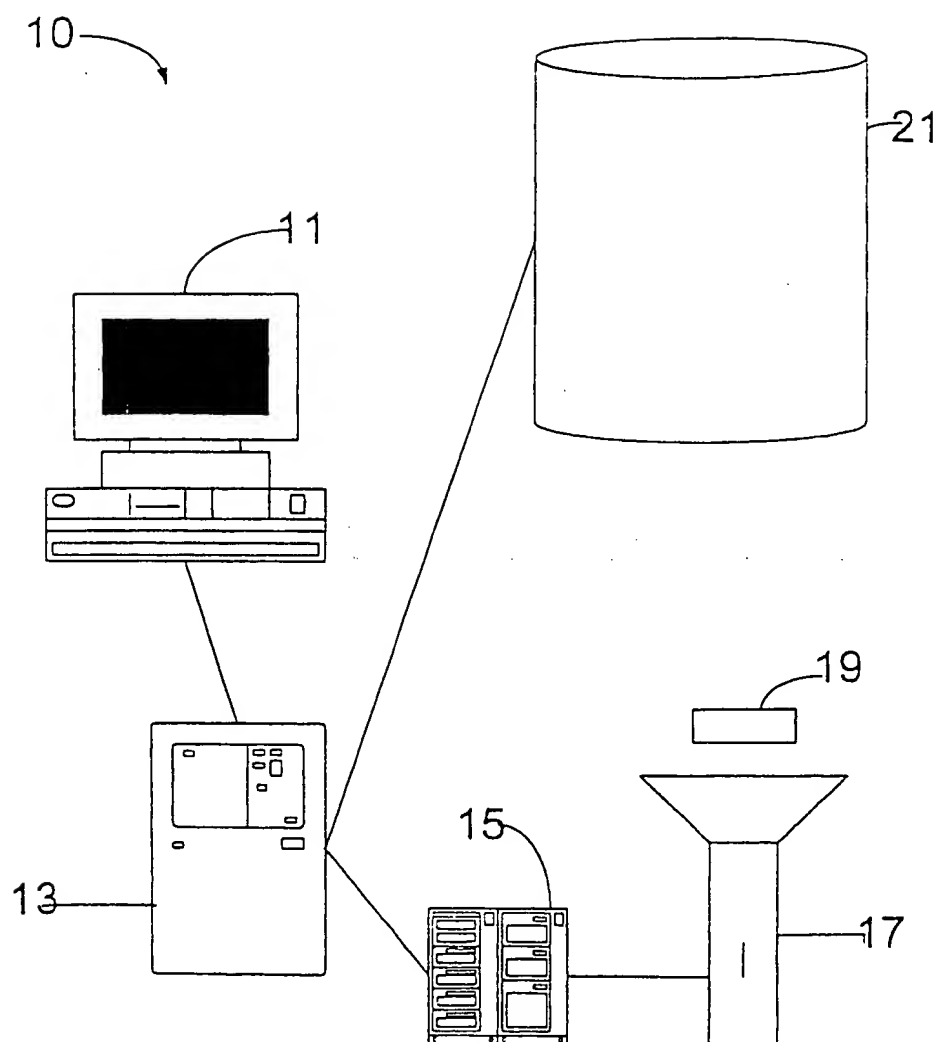


FIG. 1

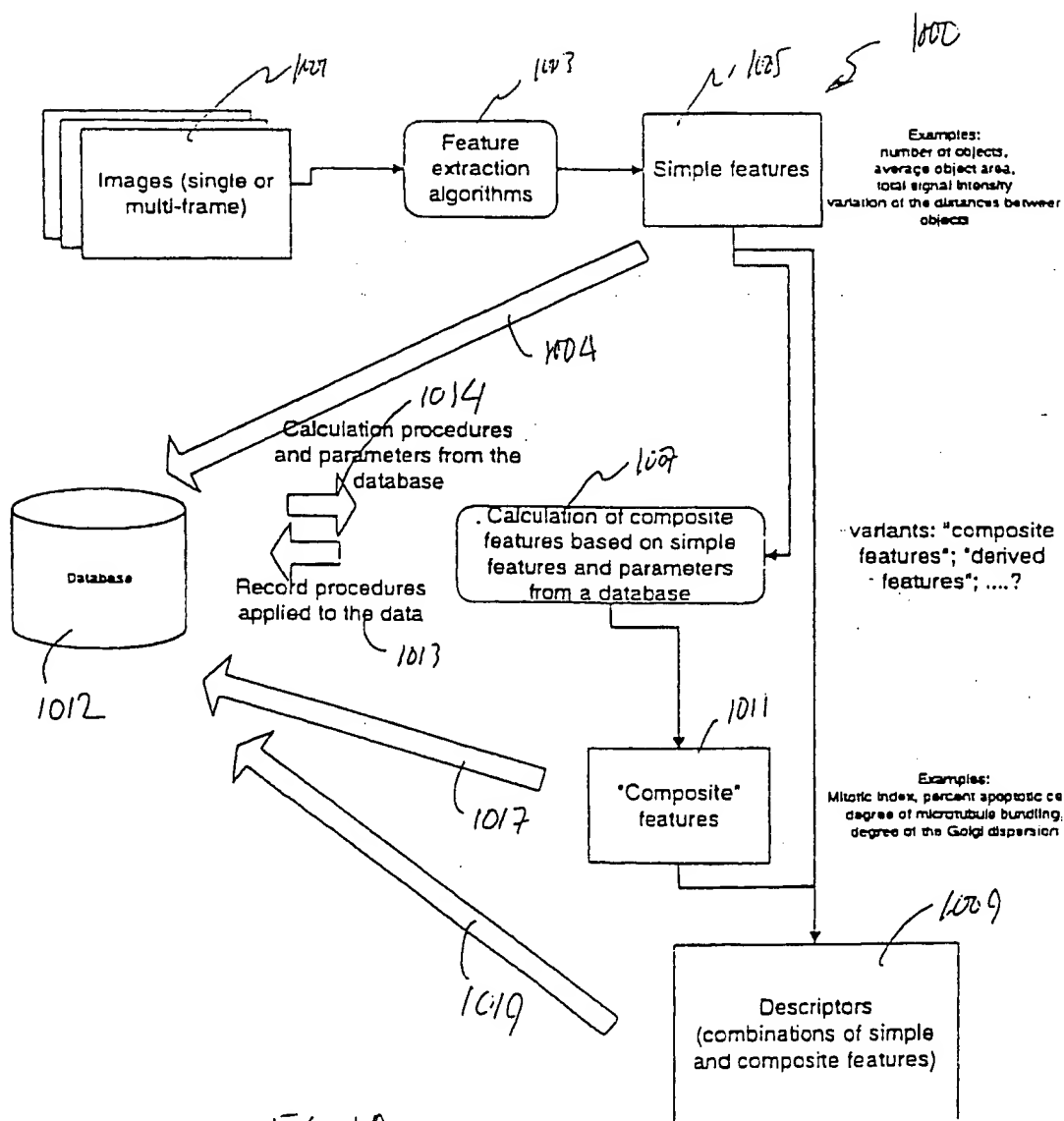


FIG. 1A

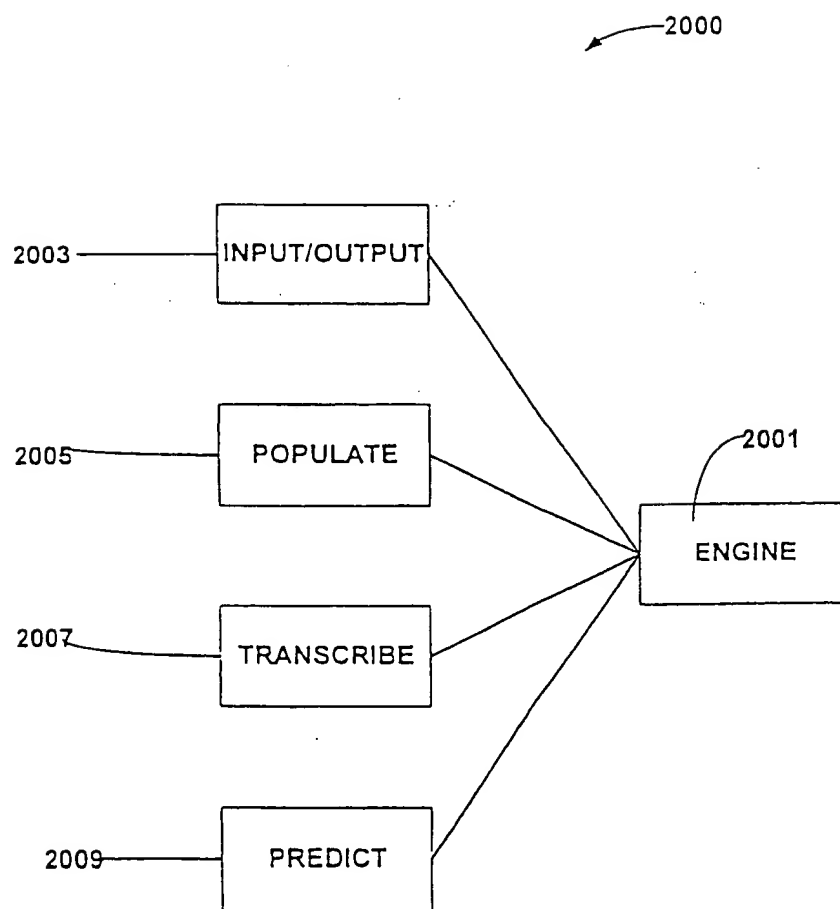


FIG. 1B

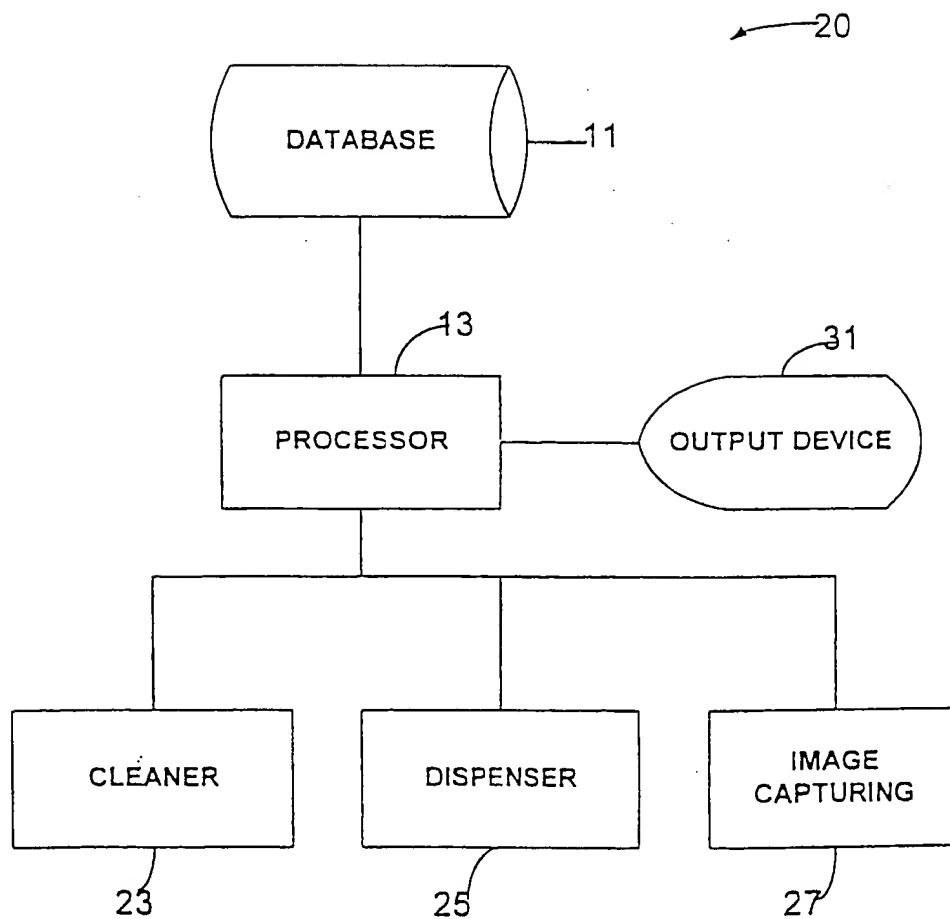


FIG. 2

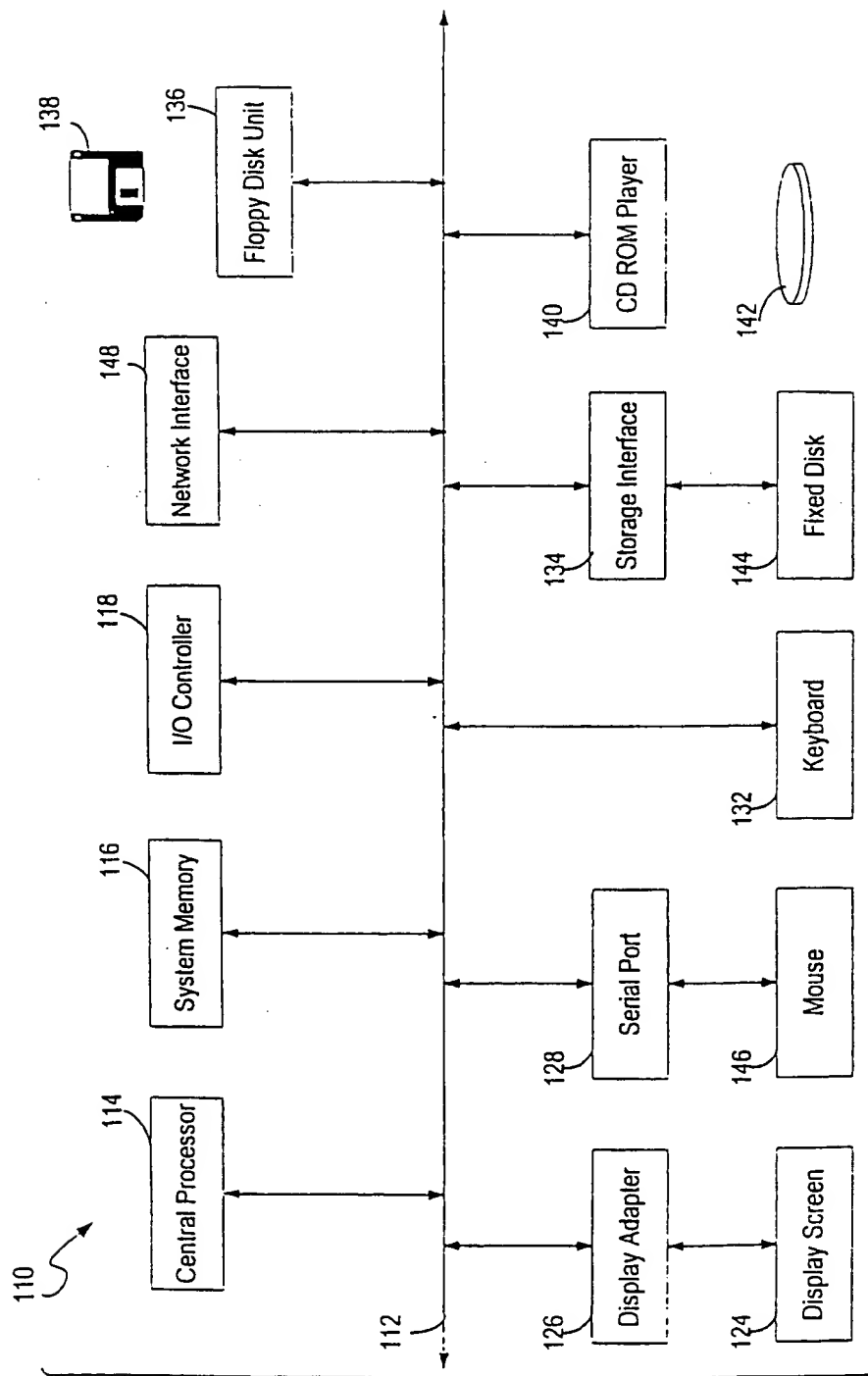


Fig 3

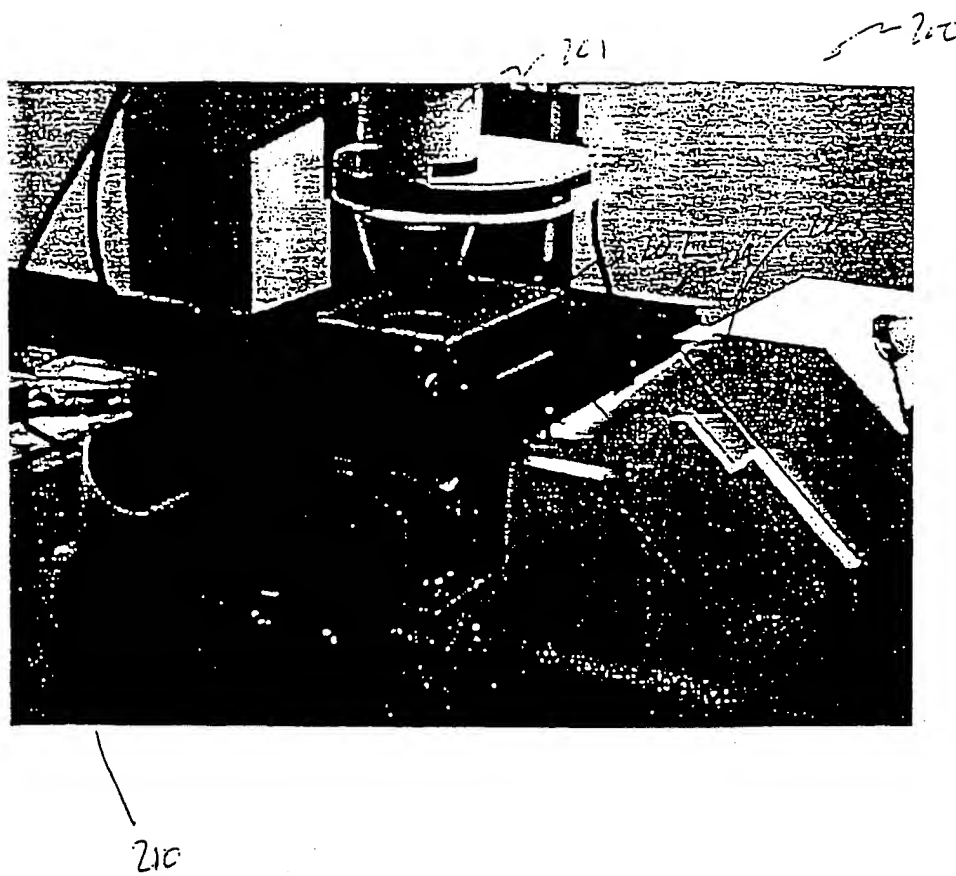


FIG. 4

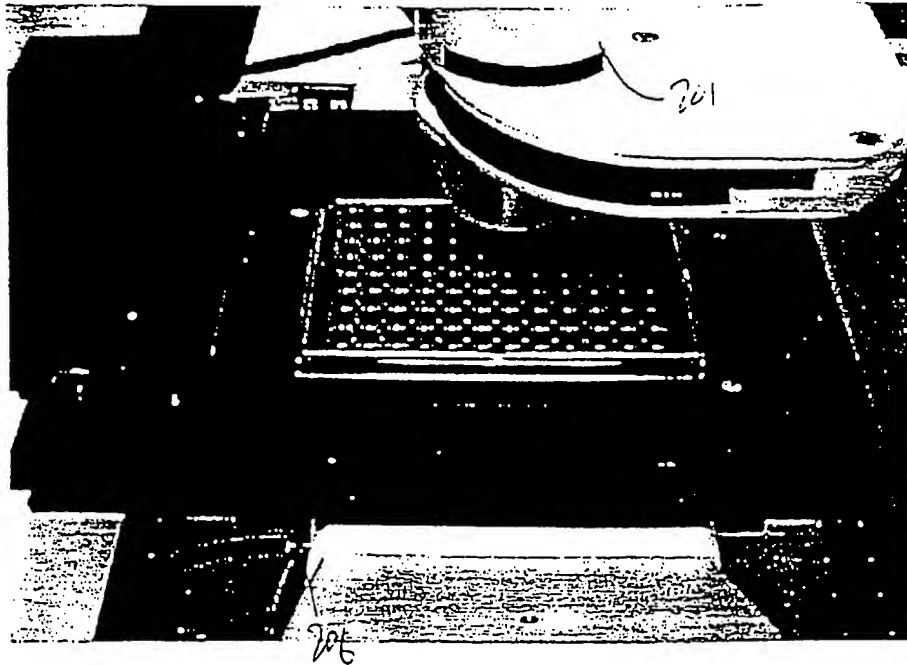


FIG 5

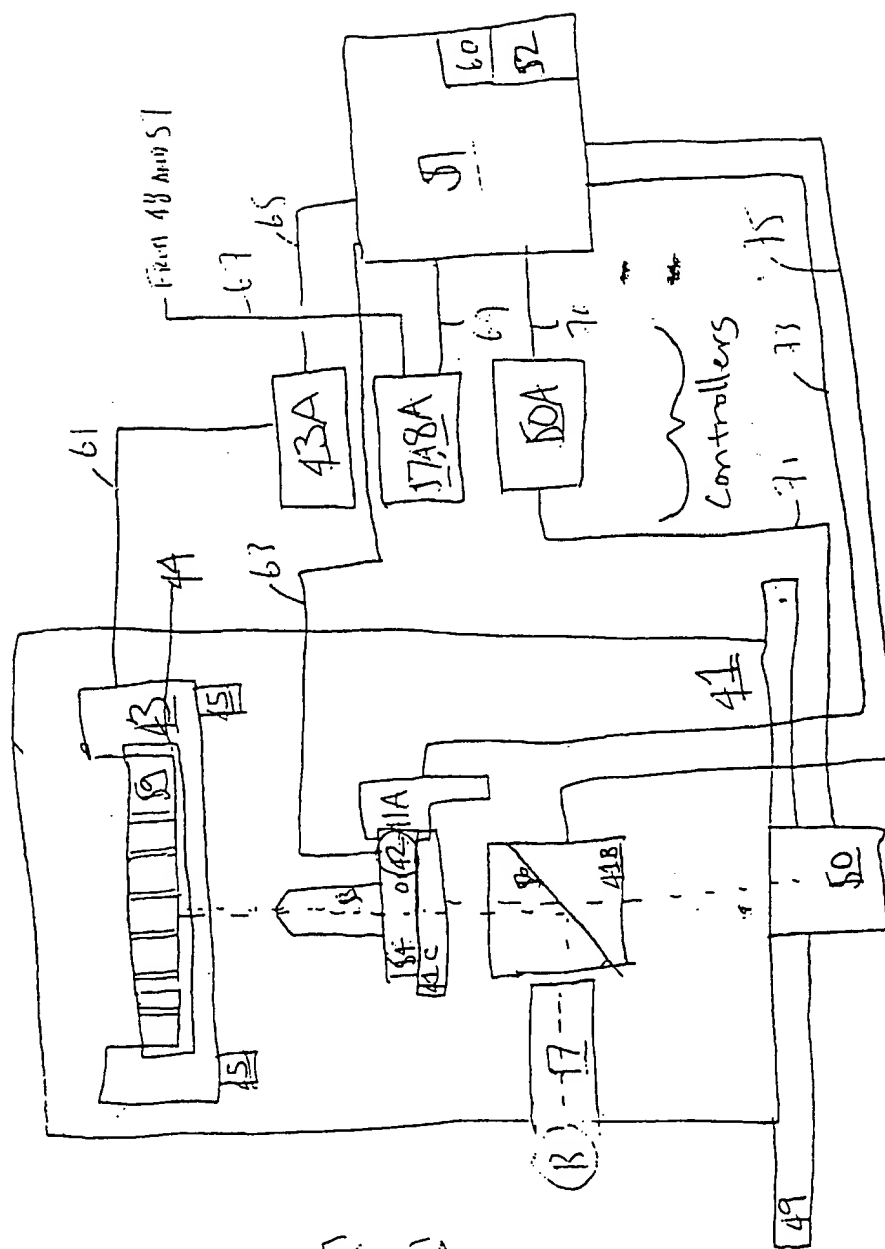


FIG. 5A

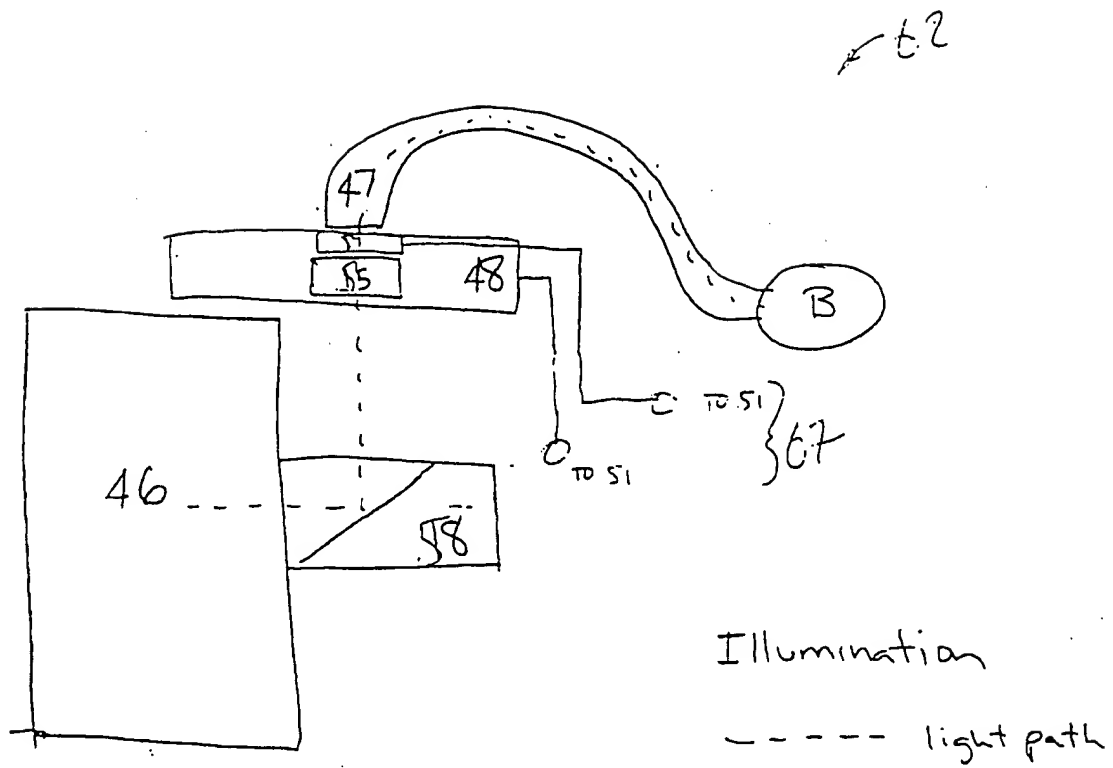


FIG. 5B

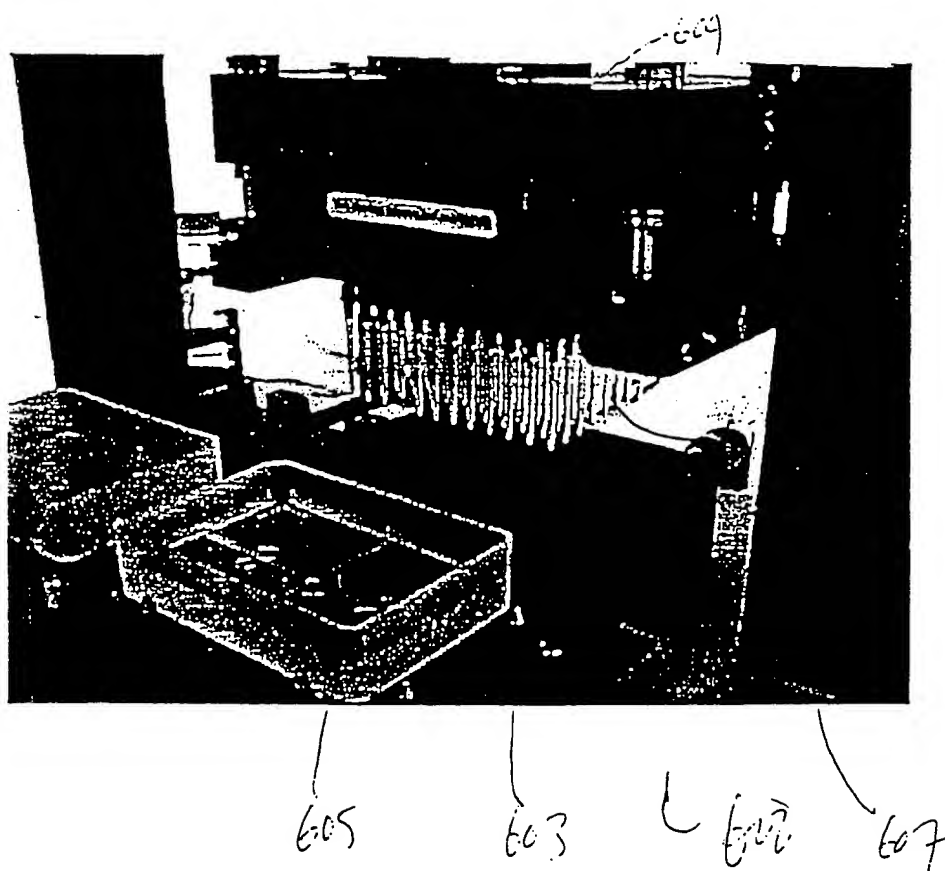


FIG. 6

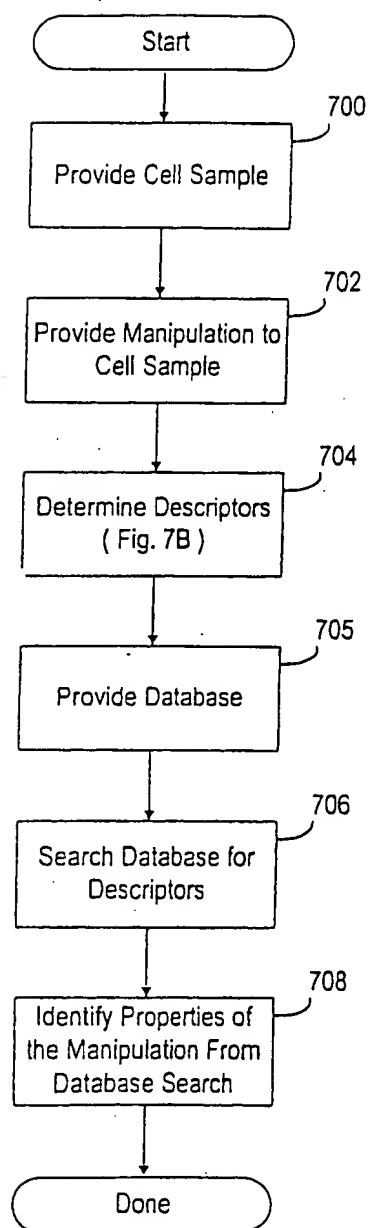
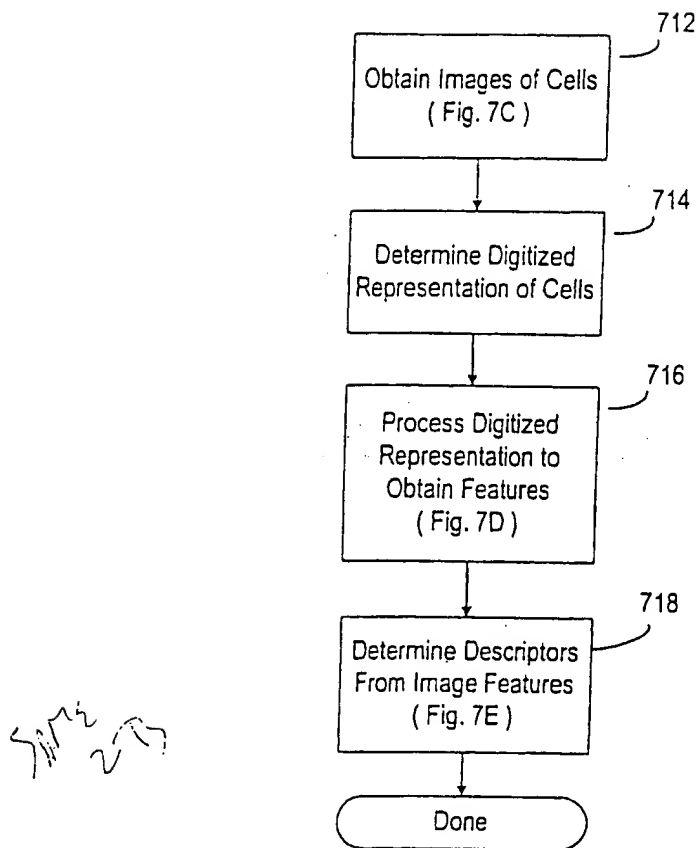


Fig. 7A



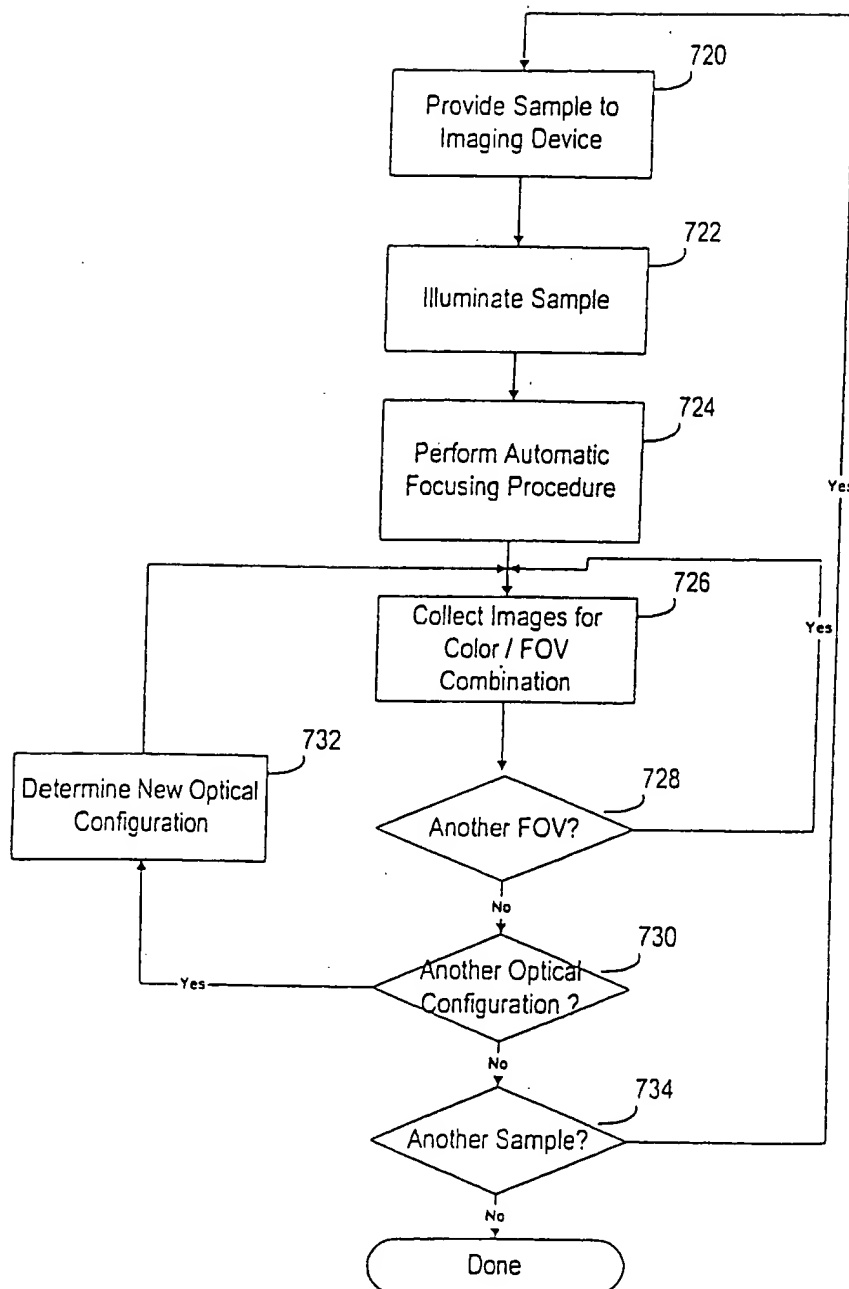


Fig. 7C
Step 714 of Fig. 7B

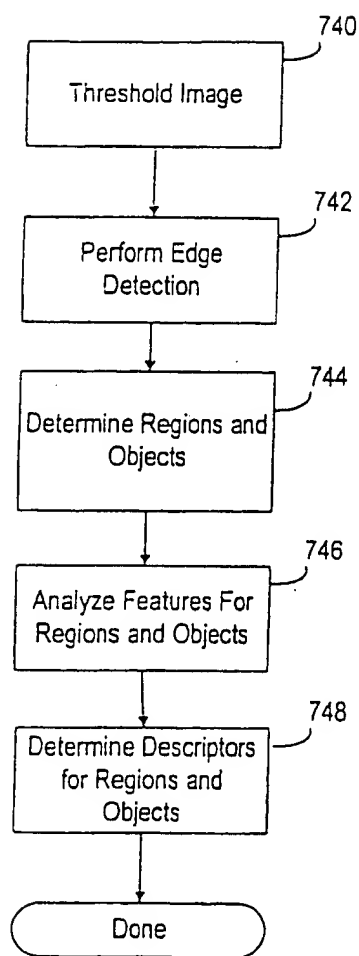


Fig. 7D
Step 716 of Fig. 7B

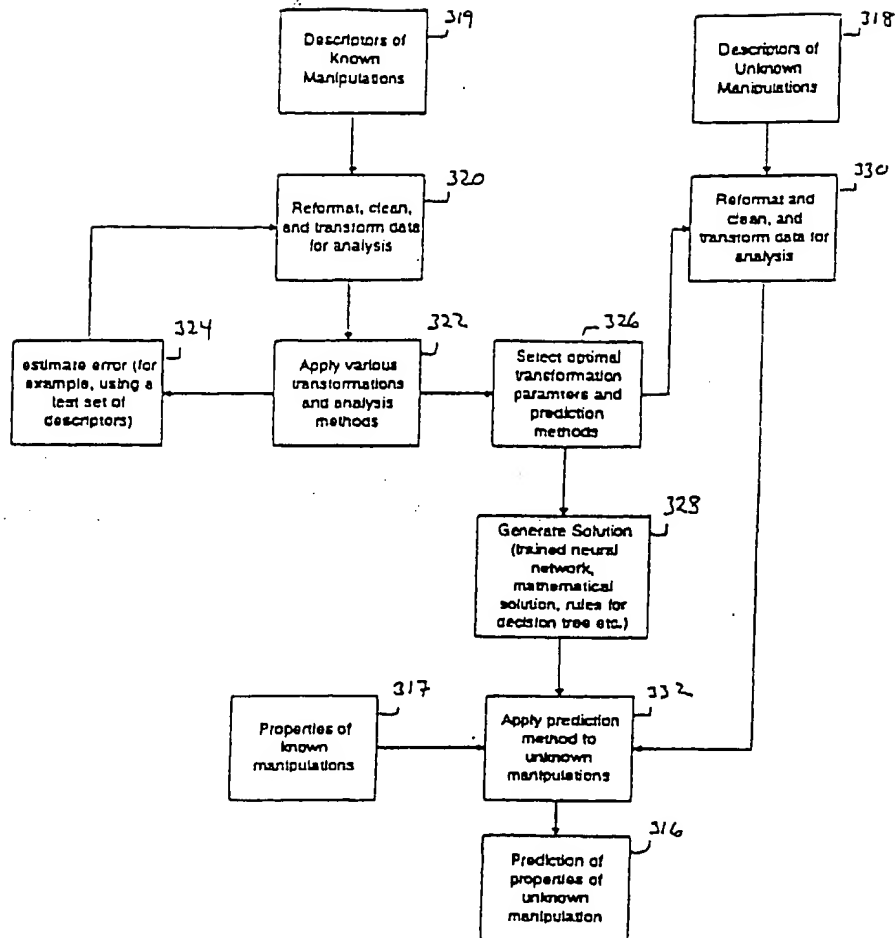


FIG. 7E

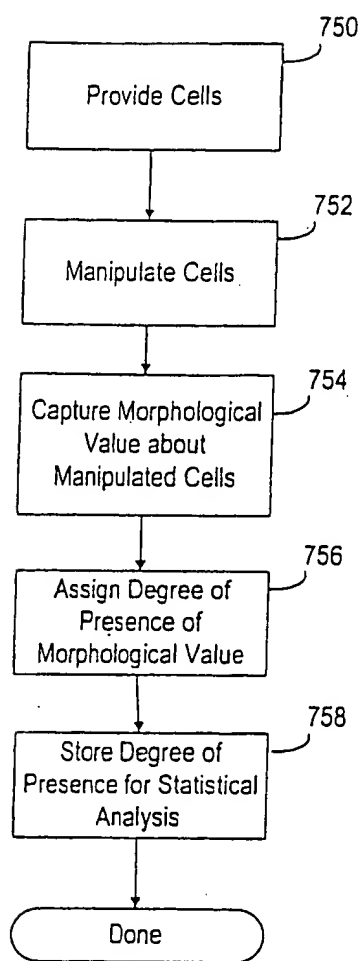


Fig. 7F

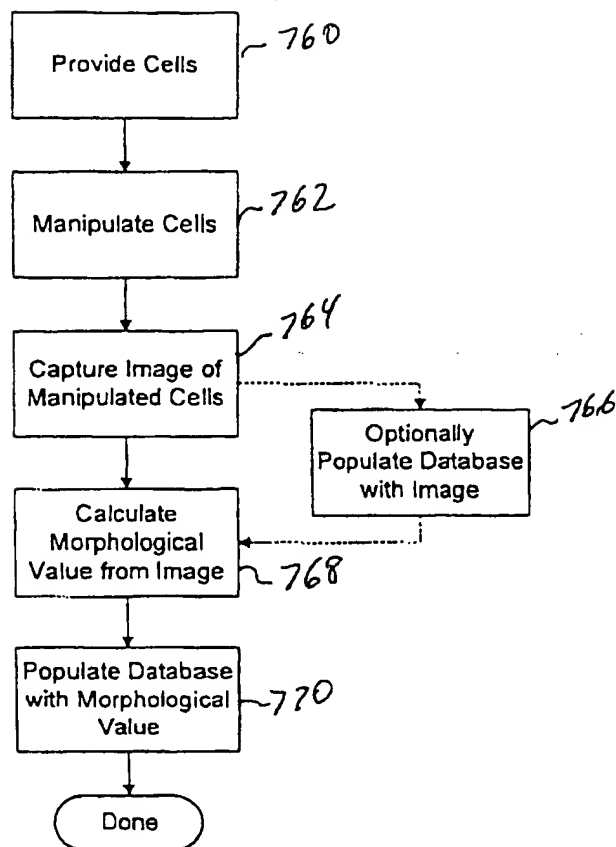


Fig 76

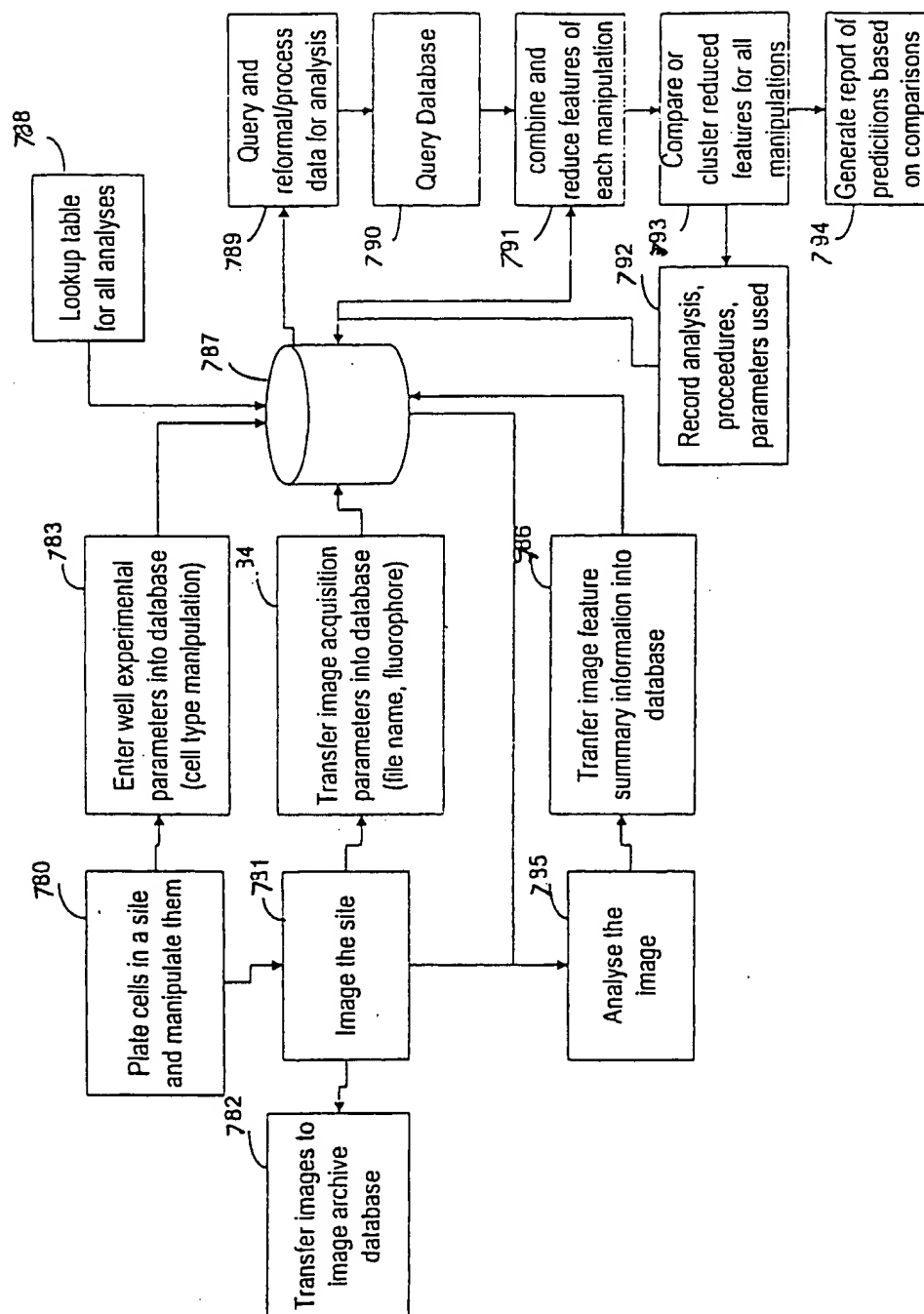


Fig.7 H

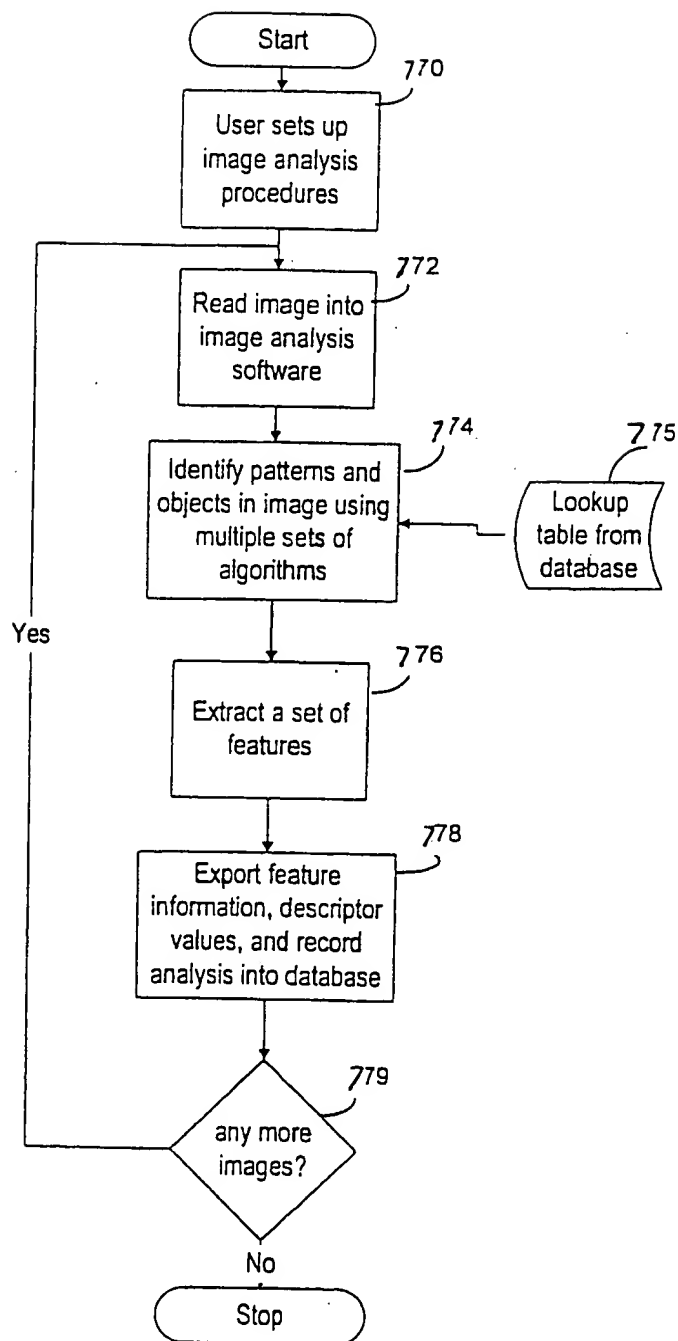
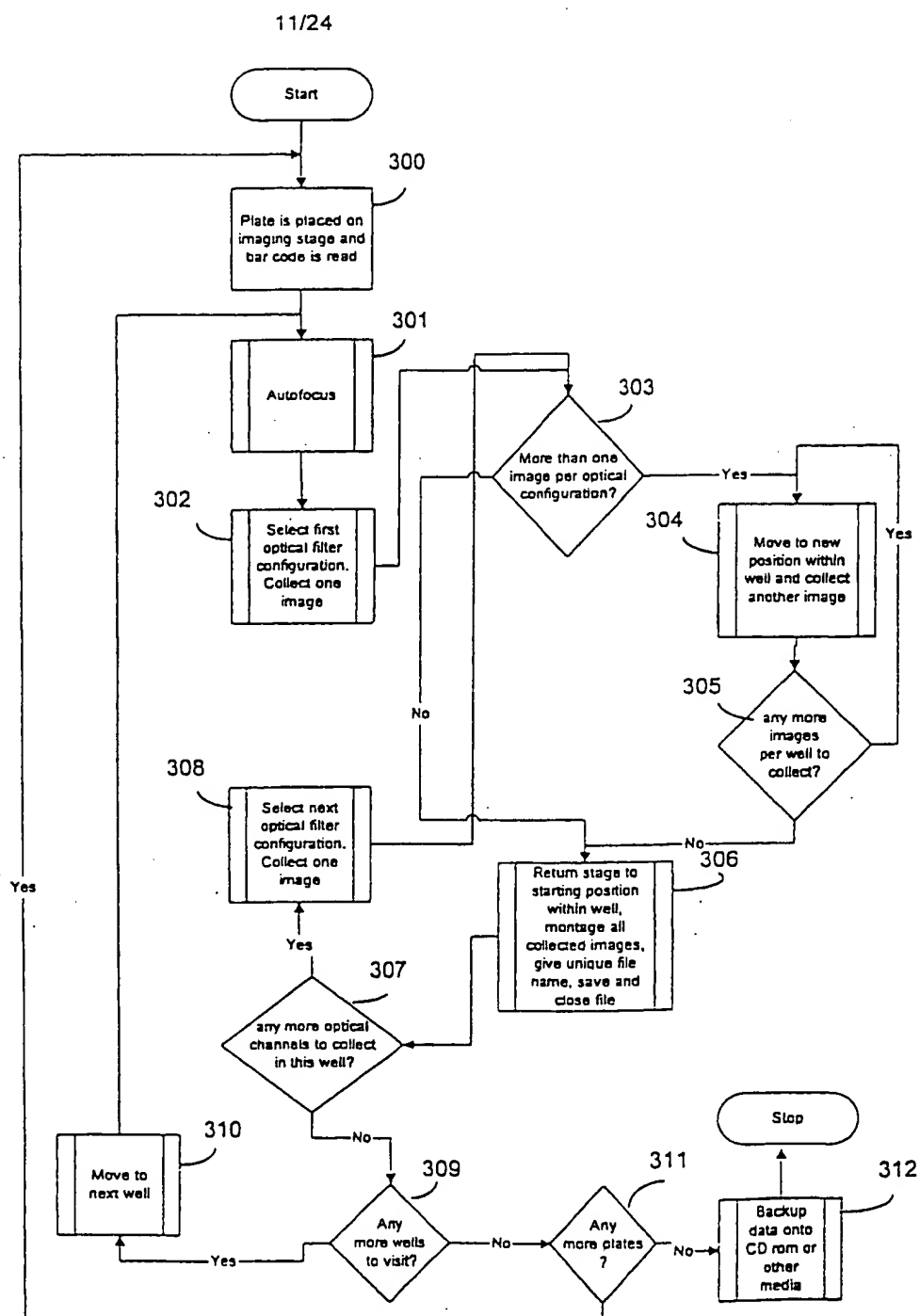


Fig. 71



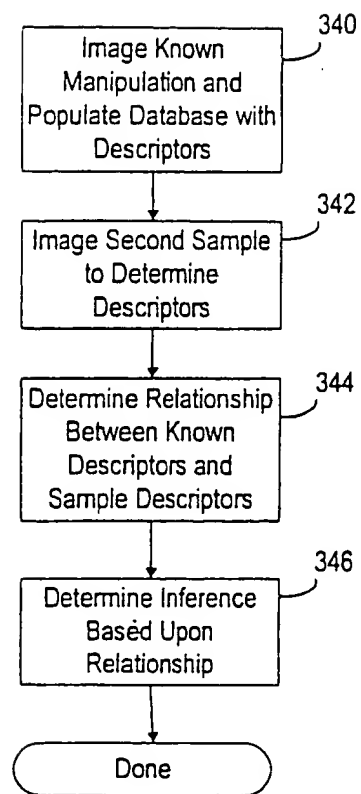


Fig. 7K

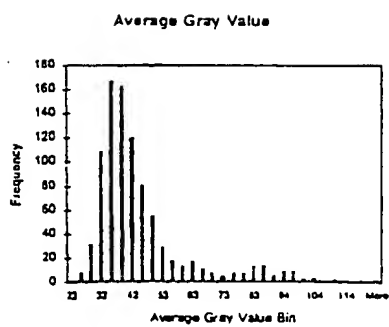


Fig. 8A

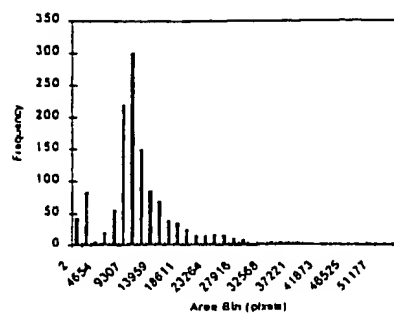


Fig. 8B

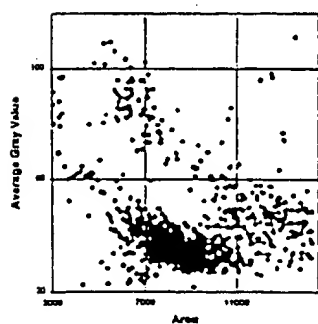


Fig. 8C

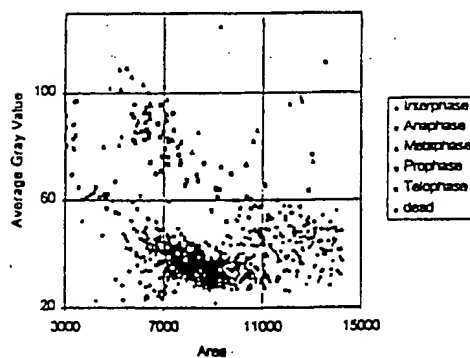


Fig. 8D

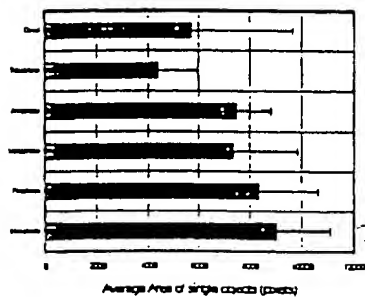


Fig. 8E

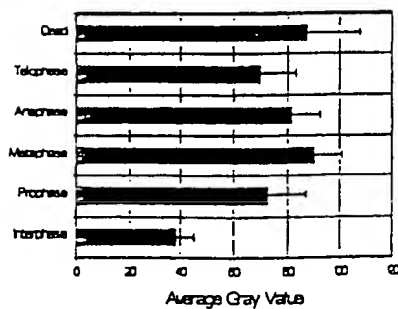


Fig. 8F

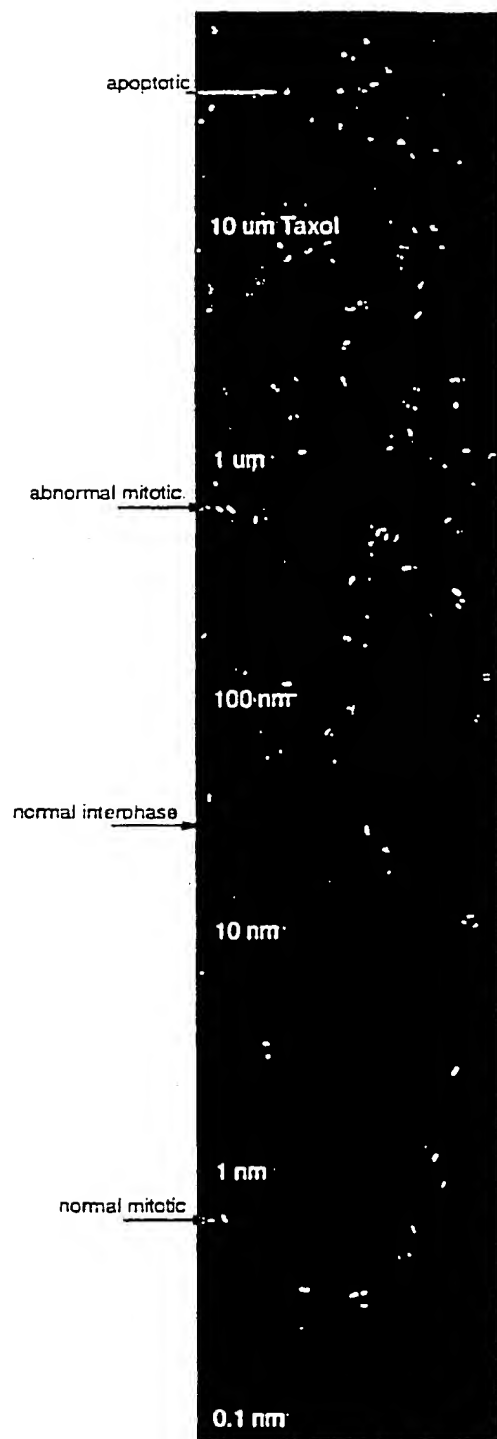


Fig. 9

MDCK cells treated with Taxol for 4.5 hours

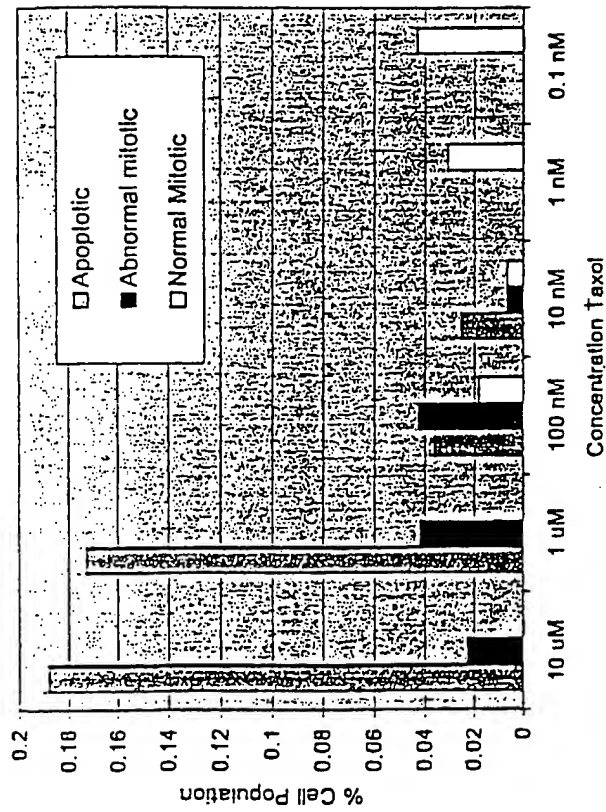


Fig. 10

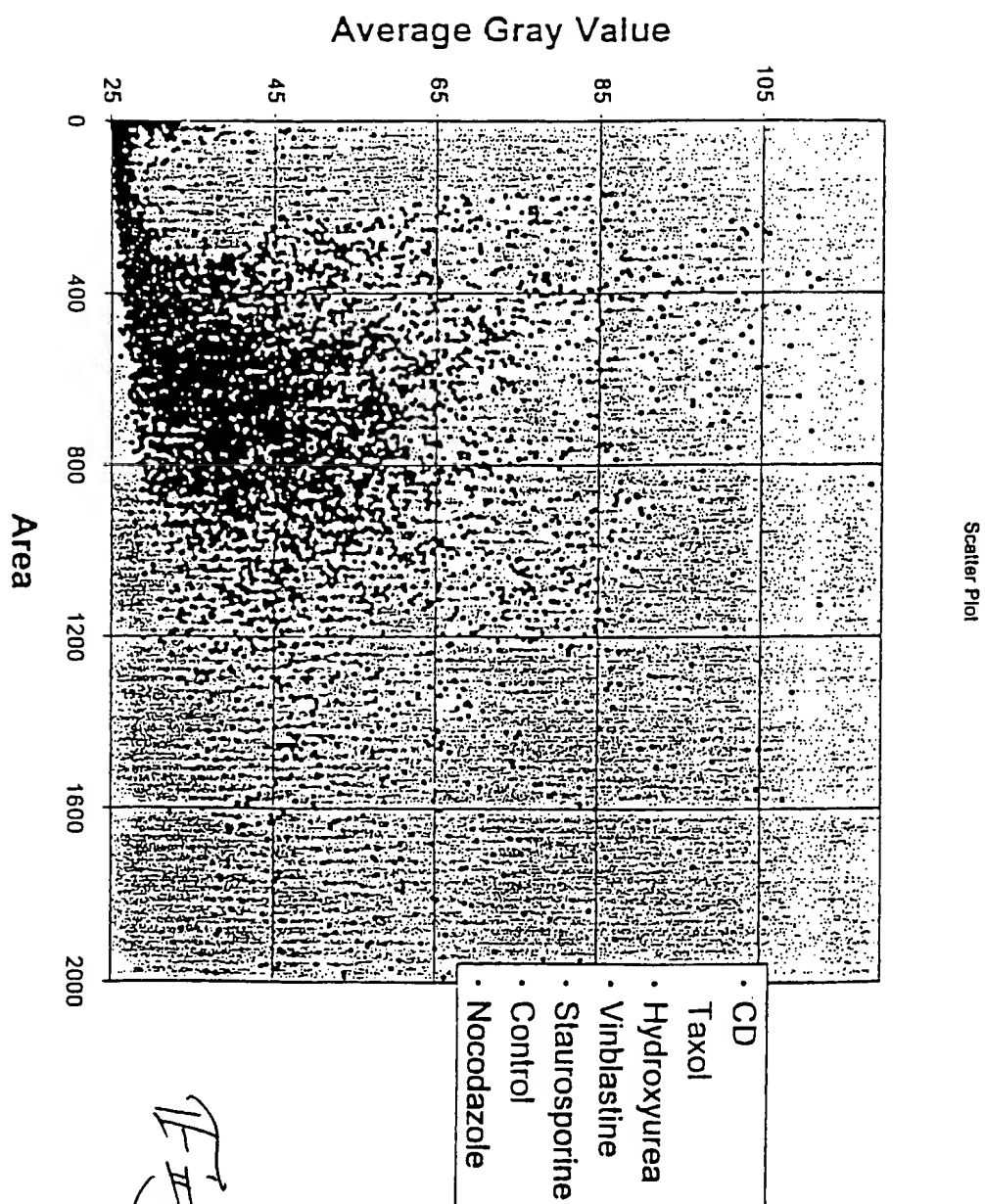
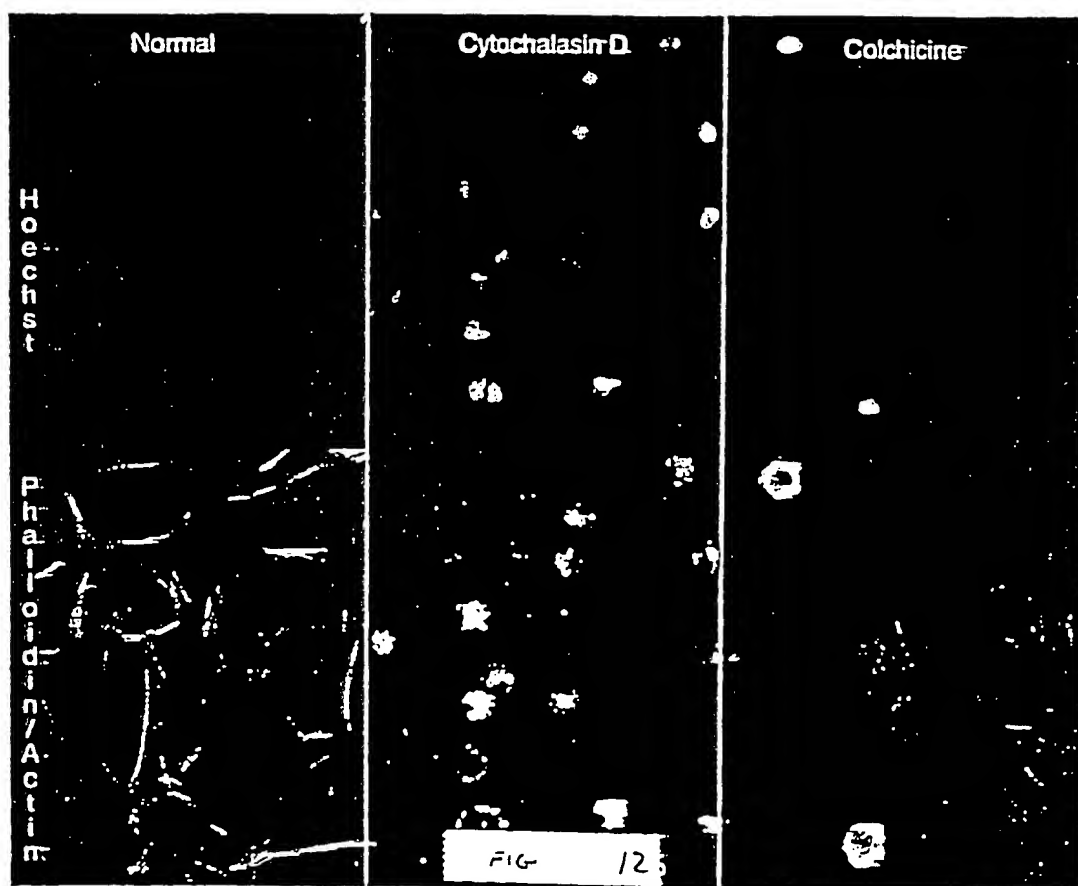


Fig 11



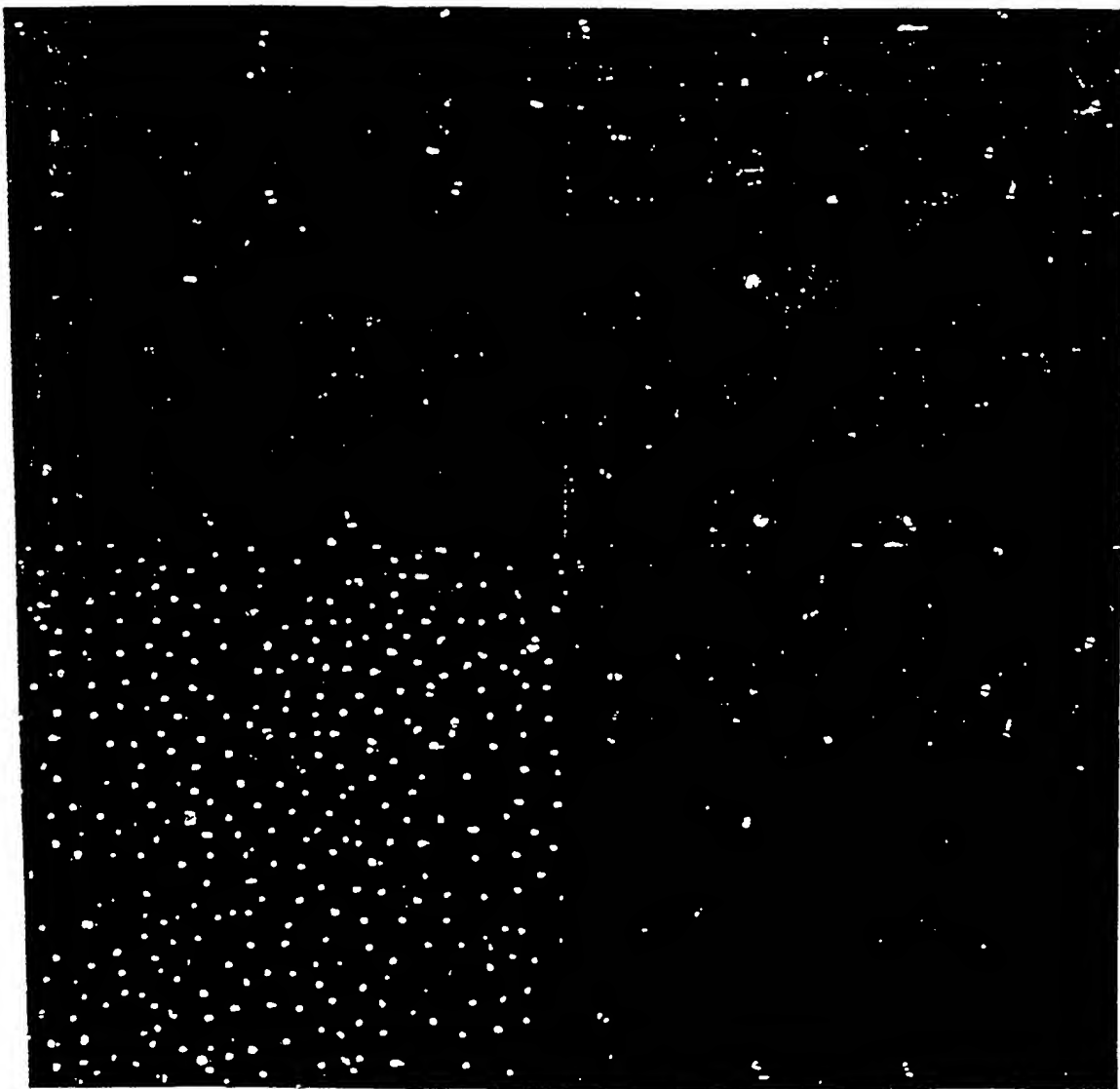


Fig 13

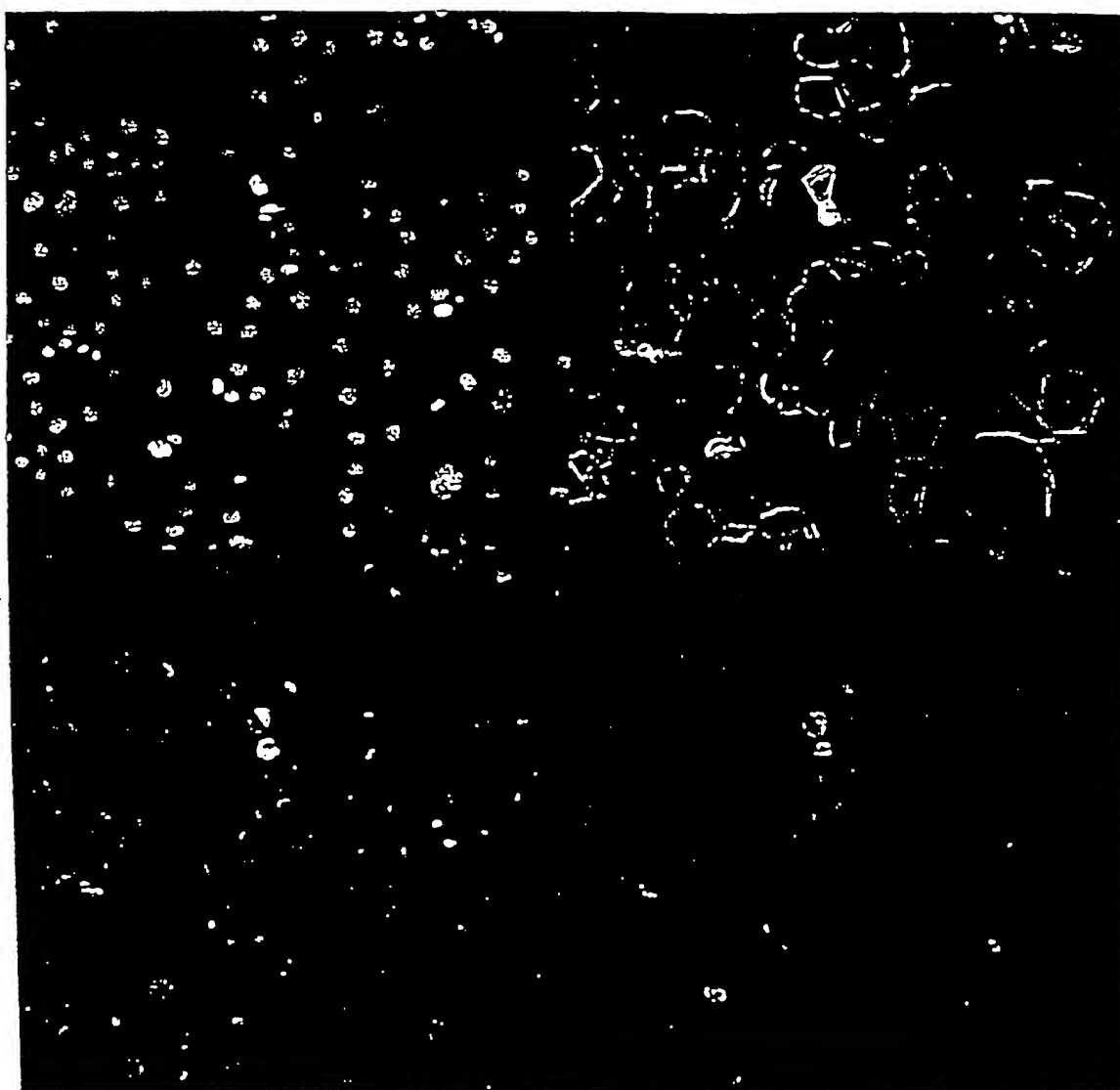


Fig 14

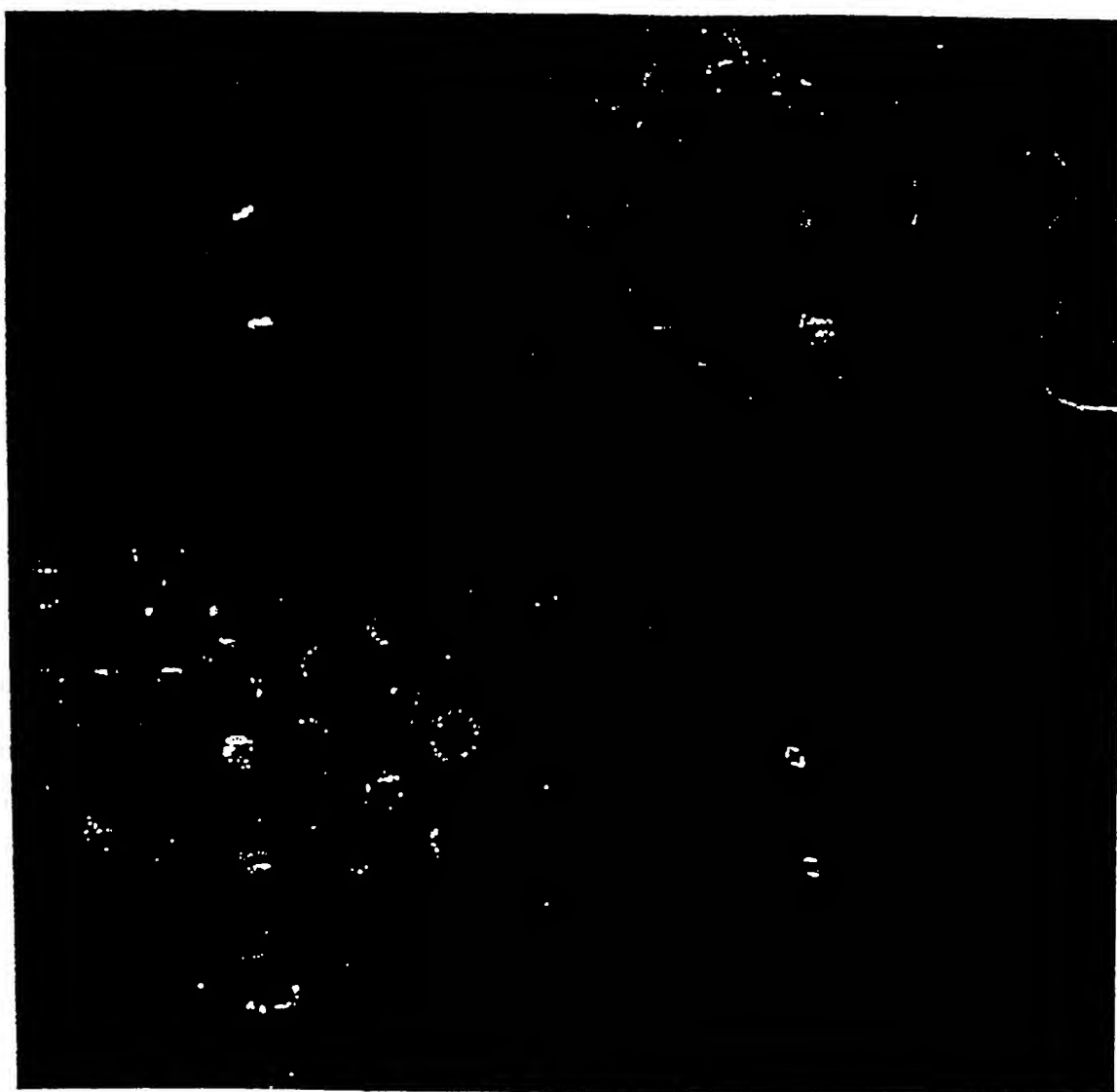


Fig 15

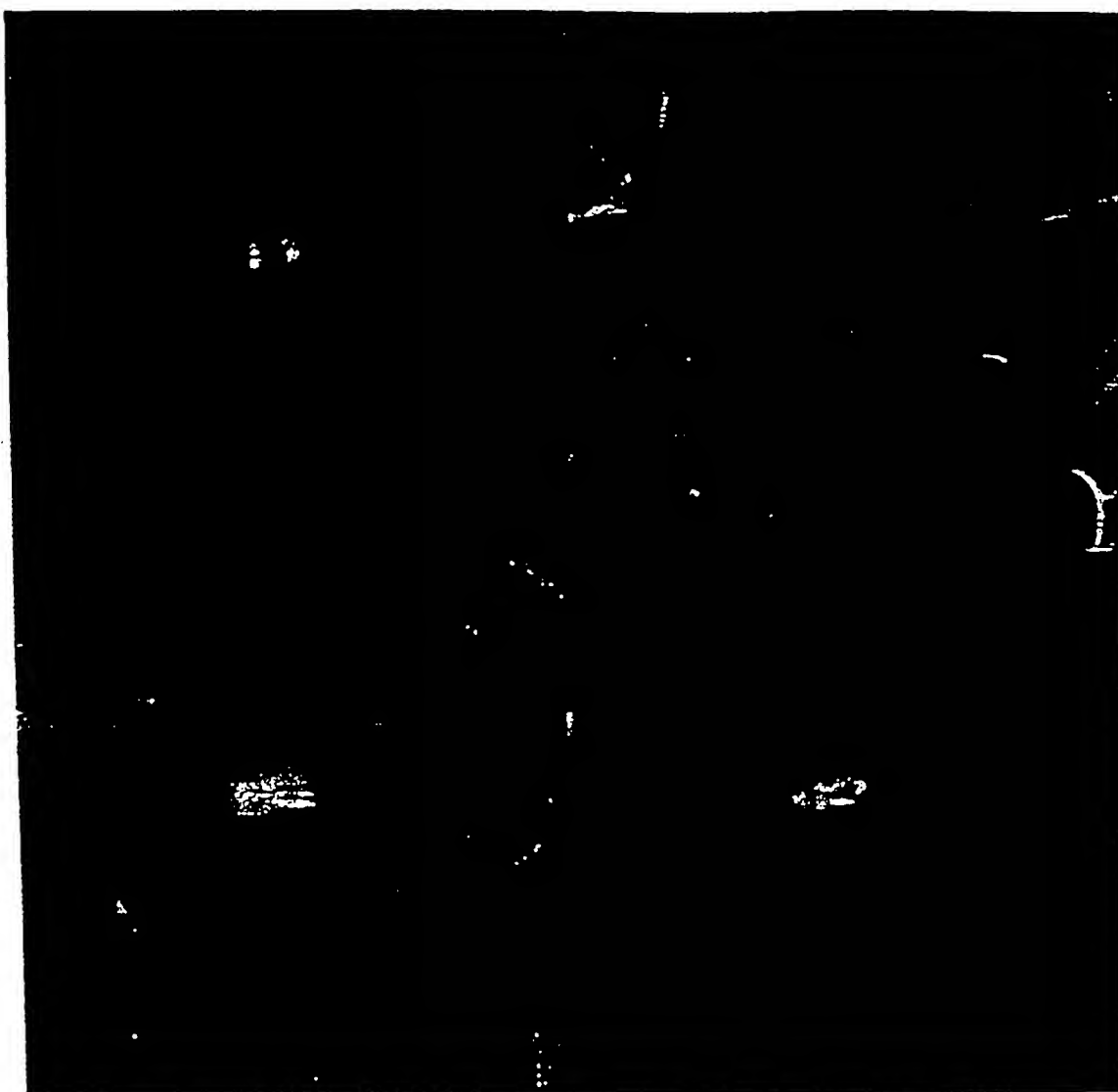


Fig 16

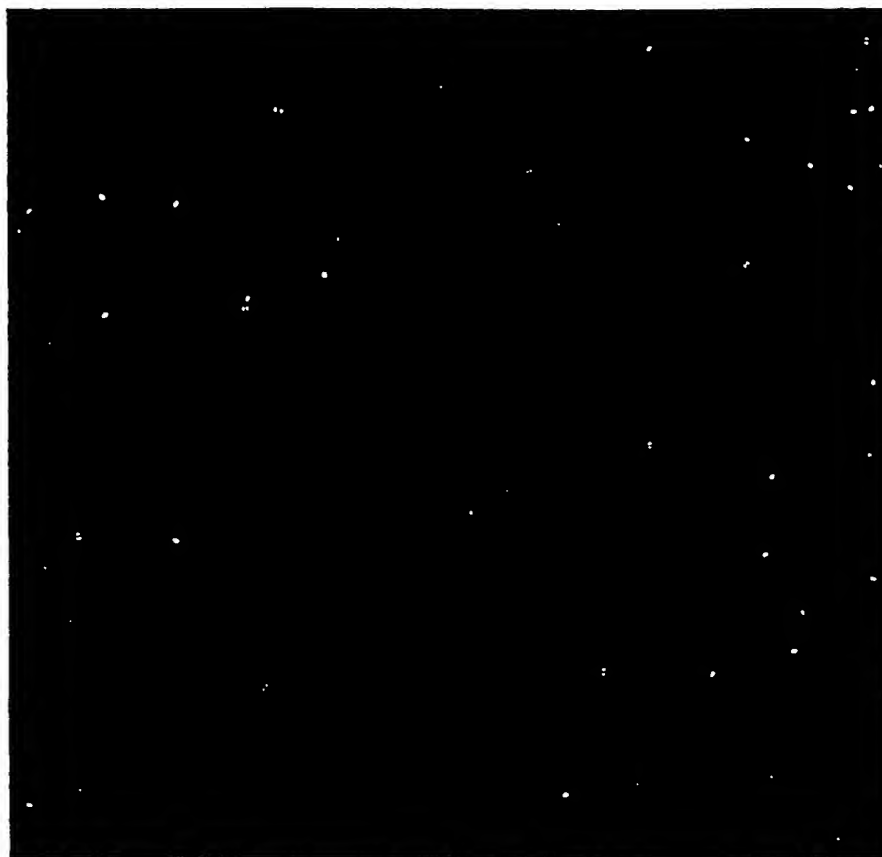


Fig 17

Conversion of morphometric parameters into nucleic acid code
and clustering of the resulting sequences using Neighbor
Joining method.

Compound:	Measurements																							
	Count	Area	Perimeter	Length	Breadth	Fiber length	Fiber breadth	Shape factor	Ell. form factor	Inner radius	Outer radius	Mean radius	Equiv. radius	Equiv. sphere vol.	Equiv. prolate vol.	Equiv. oblate vol.	Equiv. sphere surface area	Average gray value	Total gray value	Optical density	Radial dispersion	Texture Difference Moment	EFA Harmonic 2, Semi-Maj	EFA Harmonic 2, Semi-Min
Control	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	a	t	t
Taxol	a	t	t	t	t	t	t	t	a	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t
CD	c	a	a	a	t	a	t	c	a	a	a	a	a	a	a	a	a	t	a	a	a	t	a	g
Nocodazol	c	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t
Staurosporine	g	g	c	a	a	t	a	a	t	g	a	a	a	t	g	g	g	a	a	t	a	t	a	a
Vinblastine	c	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	g	t	t	t	t	t	t
Hydroxyurea	g	t	t	t	t	t	t	g	t	t	t	t	t	t	t	t	t	t	t	c	t	a	t	t

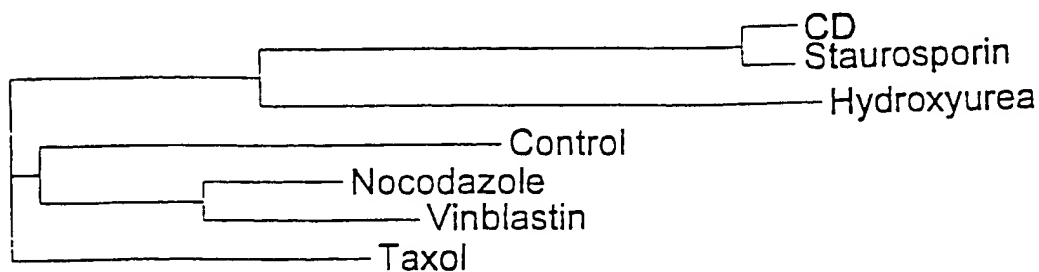


Fig 10

Conversion of morphometric parameters into amino acid codes
and clustering of the resulting sequences using Neighbor
Joining method.

	Count	Area	Perimeter	Length	Breadth	Fiber length	Fiber breadth	Shape factor	Ell. form factor	Inner radius	Outer radius	Mean radius	Equiv. radius	Equiv. sphere vol.	Equiv. prolate vol.	Equiv. oblate vol.	Equiv. sphere surface a	Average gray value	Total gray value	Optical density	Radial dispersion	Texture Difference Mo	IEFA Harmonic 2, Semi-	IEFA Harmonic 2, Semi-
Control	H	P	T	T	Z	S	D	W	F	S	T	T	T	T	C	C	P	P	M	C	T	G	T	T
Taxol	G	F	M	M	P	M	P	H	G	S	M	M	W	C	F	P	F	R	C	M	M	H	M	P
CD	F	G	G	G	M	G	M	K	A	G	G	G	G	G	G	G	H	G	G	G	M	G	V	H
Nocodazole	W	F	M	M	W	M	P	T	R	S	M	M	M	F	M	W	F	M	M	R	M	M	M	F
Staurosporine	N	V	A	G	G	M	G	Y	V	G	G	G	M	V	V	V	G	G	H	G	M	G	G	V
Vinblastine	F	W	W	M	W	W	C	W	D	S	M	W	W	M	M	M	W	M	V	E	M	M	M	F
Hydroxyurea	S	H	H	H	H	H	H	V	H	H	H	H	H	H	H	H	H	H	H	A	H	G	H	D

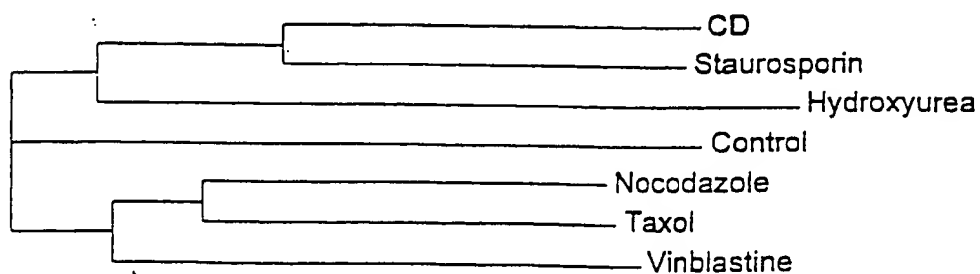


Fig 19

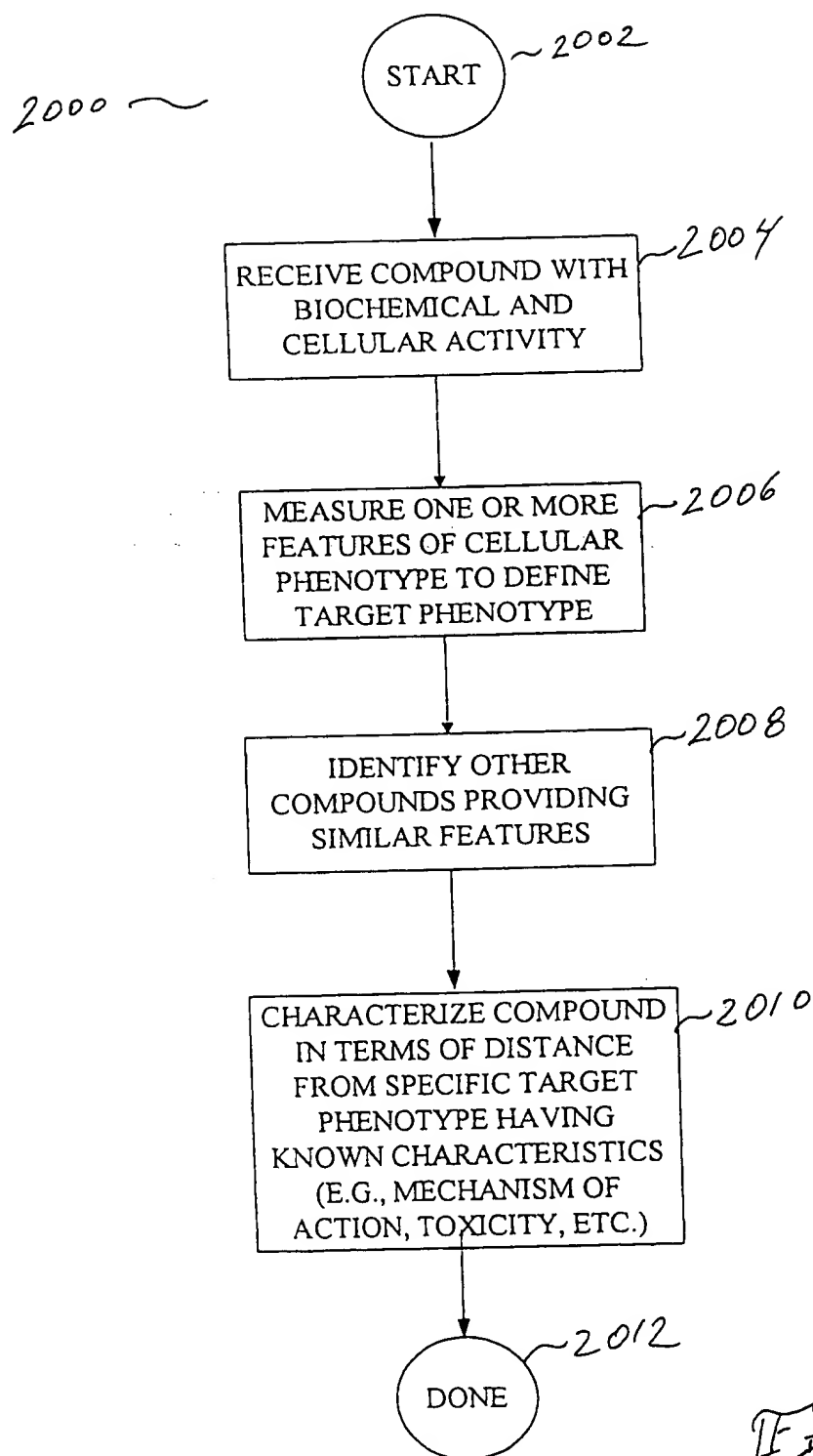


Fig 20

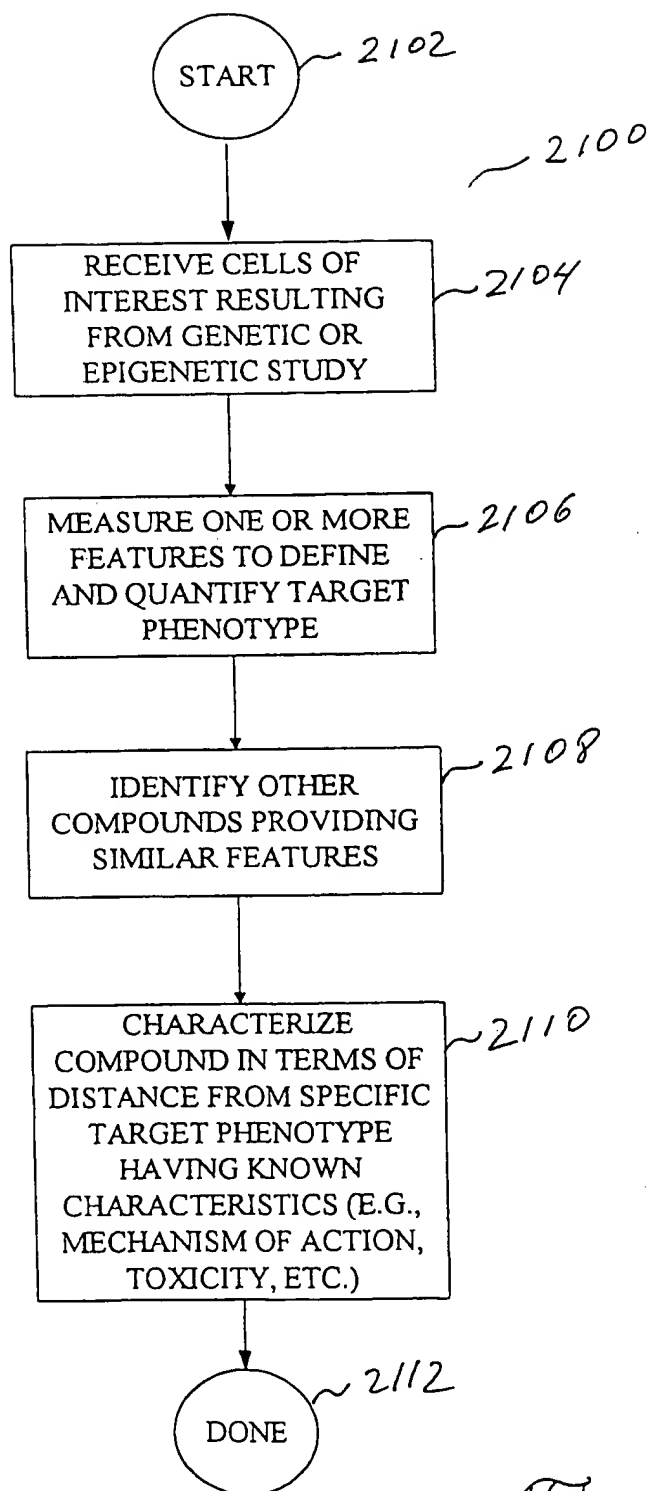


Fig 21

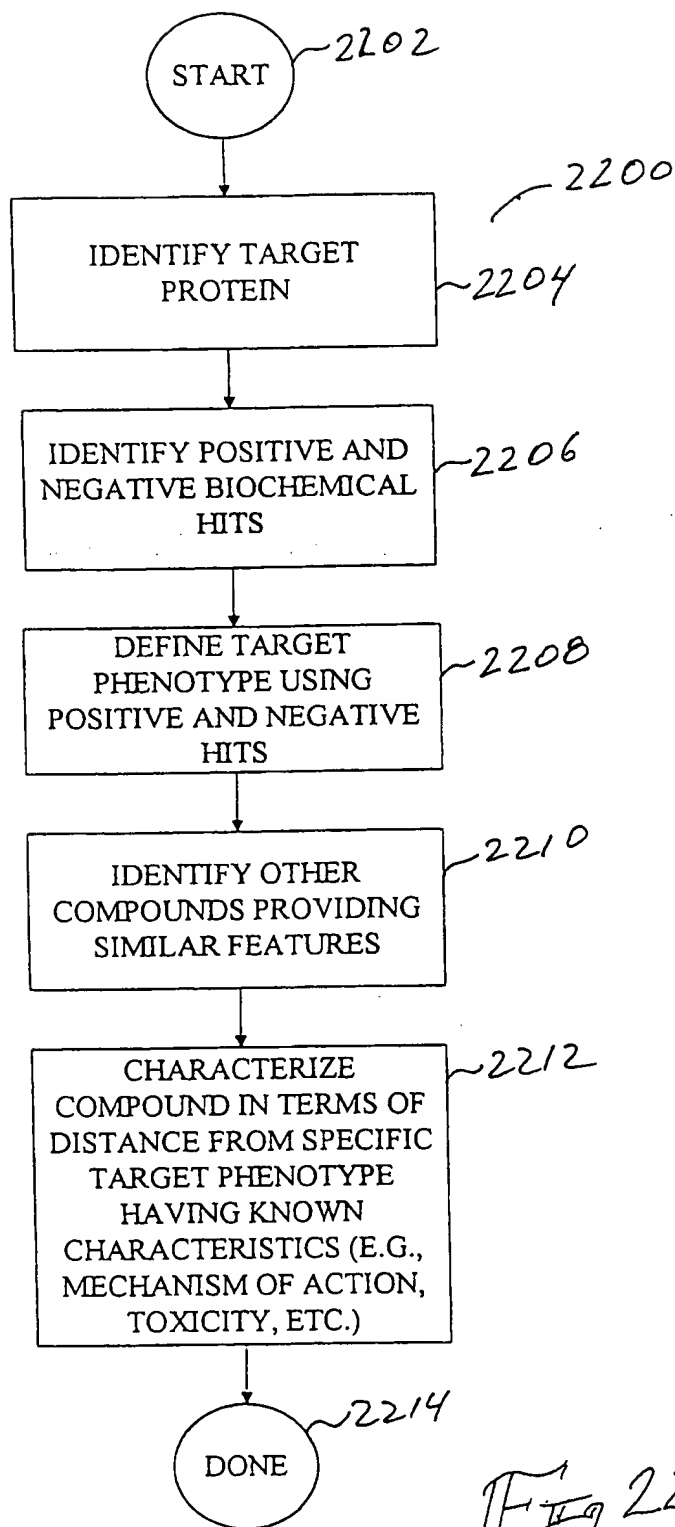


Fig 22

(19) World Intellectual Property Organization
International Bureau



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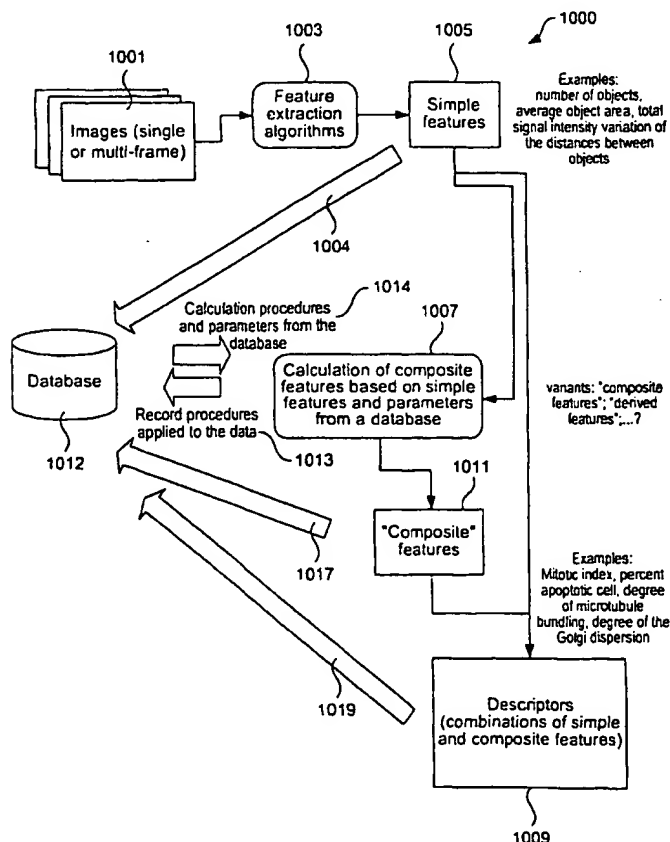
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- (74) Agent: LOUIE, Michael, L.; Beyer Weaver & Thomas, LLP, P.O. Box 130, Mountain View, CA 94042-0130 (US).
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- (71) Applicant: CYTOKINETICS, INC. [US/US]; Suite 2, 280 East Grand Avenue, South San Francisco, CA 94080 (US).

[Continued on next page]

(54) Title: METHOD AND APPARATUS FOR PREDICTIVE CELLULAR BIOINFORMATICS



(57) Abstract: Techniques for using information technology in therapeutics or drug discovery. In an exemplary embodiment, techniques for determining information about the properties of substances based upon information about structure of living or non-living cells exposed to substances are provided. A method according to the present invention enables researchers and/or scientists to identify promising candidates in the search for new and better medicines or treatments using, for example, a cellular informatics database. The present invention further teaches a system for acquiring knowledge from cellular information. The system has a database 1012 comprising a database management module ("DBMS"). The system also has a variety of modules, including a population module coupled to the DBMS for categorizing and storing a plurality of features (e.g., cell size, distance between cells, cell population, cell type) from an image acquisition device into the database. The system has a translation module coupled to the DBMS for defining a descriptor from a set of selected features from the plurality of features. In a specific embodiment, the descriptor is for a known or unknown compound, e.g., drug. A prediction module is coupled to the DBMS for selecting one of a plurality of descriptors from known and unknown compounds from the database based upon a selected descriptor from a selected compound. The selected compound may be one that is useful for treatment of human beings or the like.



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International Application No

PCT/US 00/13154

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According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 G06F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 98 38490 A (BIODX INC ;DUNLAY R TERRY (US); GOUGH ALBERT H (US); GIULIANO KENN) 3 September 1998 (1998-09-03) cited in the application	1-6, 24-27
Y	page 1; claims 1-43	7-23
X	WO 98 45704 A (TULLIN SOEREN ;KASPER ALMHOLT (DK); NOVONORDISK AS (DK); SCUDDER K) 15 October 1998 (1998-10-15) abstract; claims 1-3,22,73,80,81,86	1-6, 24-27

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040 Tx 31 651 epo.nl

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INTERNATIONAL SEARCH REPORT

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PCT/US 00/13154

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	<p>MONTIRONI R ET AL: "COMPUTED CELL CYCLE AND DNA HISTOGRAM ANALYSES IN IMAGE CYTOMETRY IN BREAST CANCER"</p> <p>JOURNAL OF CLINICAL PATHOLOGY, GB, LONDON, vol. 46, no. 9, 1 September 1993 (1993-09-01), pages 795-800, XP000644549</p> <p>ISSN: 0021-9746</p> <p>abstract</p> <p style="text-align: center;">---</p>	7-13
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A	<p>GIULIANO K A ET AL: "Fluorescent-protein biosensors: new tools for drug discovery"</p> <p>TRENDS IN BIOTECHNOLOGY, GB, ELSEVIER PUBLICATIONS, CAMBRIDGE,</p> <p>vol. 16, no. 3, 1 March 1998 (1998-03-01), pages 135-140, XP004108592</p> <p>ISSN: 0167-7799</p> <p>page 139, left-hand column, paragraph 4</p> <p>-right-hand column, paragraph 3</p> <p style="text-align: center;">-----</p>	1-27

INTERNATIONAL SEARCH REPORT

Information on patent family members

Intel onal Application No

PCT/US 00/13154

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WO 0050872	A	31-08-2000	NONE	

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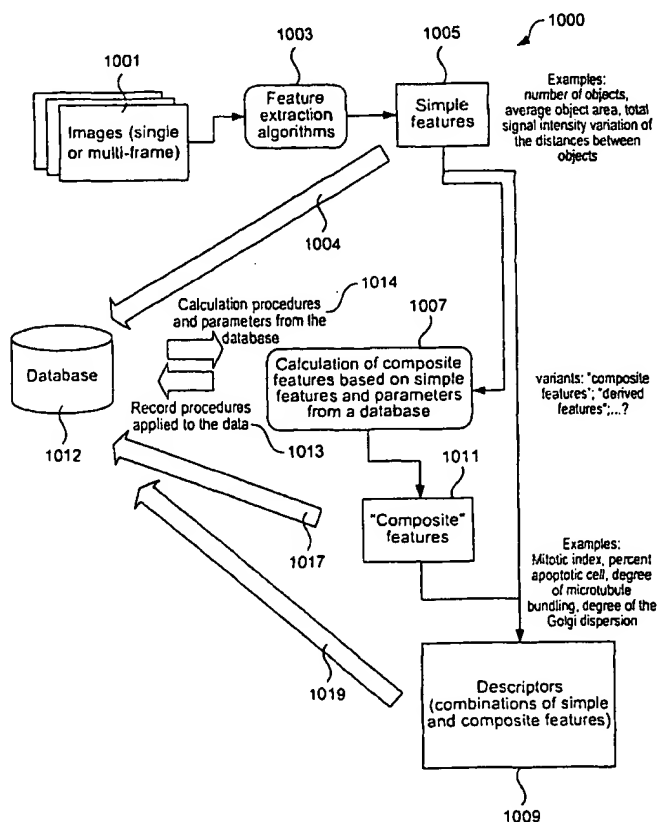
Not furnished 26 April 2000 (26.04.2000) US

(72) Inventors: **SABRY, James, H.**; 52 Buena Vista Terrace, San Francisco, CA 94117 (US). **ADAMS, Cynthia, L.**; 615 Georgia Avenue, Palo Alto, CA 94306 (US). **VAISBERG, Eugeni, A.**; 647 Pegasus Lane, Foster City, CA 94404 (US). **CROMPTON, Anne, M.**; 2 Bellair Place, San Francisco, CA 94133 (US). **BLUM, Robert, I.**; 17 Shoreview Avenue, San Francisco, CA 94121 (US). **OESTREICHER, Donald, R.**; 904 Old Town Court, Cupertino, CA 95014-4024 (US). **SIGAL, Nolan, H.**; 941 Berry Avenue, Los Altos, CA 94024 (US).

(74) Agent: **LOUIE, Michael, L.**; Beyer Weaver & Thomas, LLP, P.O. Box 130, Mountain View, CA 94042-0130 (US).

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[Continued on next page]

(54) Title: **METHOD AND APPARATUS FOR PREDICTIVE CELLULAR BIOINFORMATICS**

(57) Abstract: Techniques for using information technology in therapeutics or drug discovery. In an exemplary embodiment, techniques for determining information about the properties of substances based upon information about structure of living or non-living cells exposed to substances are provided. A method according to the present invention enables researchers and/or scientists to identify promising candidates in the search for new and better medicines or treatments using, for example, a cellular informatics database. The present invention further teaches a system for acquiring knowledge from cellular information. The system has a database 1012 comprising a database management module ("DBMS"). The system also has a variety of modules, including a population module coupled to the DBMS for categorizing and storing a plurality of features (e.g., cell size, distance between cells, cell population, cell type) from an image acquisition device into the database. The system has a translation module coupled to the DBMS for defining a descriptor from a set of selected features from the plurality of features. In a specific embodiment, the descriptor is for a known or unknown compound, e.g., drug. A prediction module is coupled to the DBMS for selecting one of a plurality of descriptors from known and unknown compounds from the database based upon a selected descriptor from a selected compound. The selected compound may be one that is useful for treatment of human beings or the like.

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(AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU,
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PATENT APPLICATION
METHOD AND APPARATUS FOR
PREDICTIVE CELLULAR BIOINFORMATICS
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10 computer codes, which may be used to implement aspects of the present invention. Assignee of the present invention reserves all rights with respect to these codes and provides notice herein. Notice is hereby given © Cytokinetics, Inc. 1999.

BACKGROUND OF THE INVENTION

 The present invention provides techniques for information
15 management using a database platform. More particularly, the present invention provides a system including computer code that couples to a database device. The system provides for image capturing of living, dead, or fixed cells or cell fractions used to identify information about substances used on the cells or information about the cells themselves. Accordingly, the present invention can enable researchers and
20 scientists to identify promising candidates in the search for new and better medicines, for example, in drug discovery and development. The principles enumerated herein may, with equal facility, be applied to other applications, including but not limited to use in environmental applications such as determining chemical toxicities and other non-pharmaceutical toxicology uses.

25 For a long time, researchers in the pharmaceutical field have sought for better ways of searching for substances possessing properties that make them suitable as medicines. In the early days, researchers generally relied upon extracts from plants, dyes, and microbiological extracts for such substances. Examples of such substances include the pain reliever aspirin, the anti-cancer drug paclitaxel (brand
30 name TaxolTM), and the heart medication called digoxin. The number of useful medicines has generally been limited.

Purified substances having desirable bio-active properties are also often difficult to discover. Advances in traditional organic chemistry and more recently the rapid chemical synthesis methods often referred to as combinatorial chemistry have increased the number of compounds that researchers test for biological activity. Originally, substances were often initially tested on animals or humans to determine their biological activity. While results from such tests may identify a good drug candidate, they are often time consuming and costly, thus a limited number of substances can be tested. Therefore, pharmaceutical companies have turned to testing their ever-increasing libraries of substances against isolated proteins (drug targets) in biochemical assays that can be carried out at high throughput and low cost. It should be noted that the substances need to be tested in numerous protein tests, each customized for a particular drug target. Therefore, although each protein test may be run at a high-throughput, the design of multiple protein tests can be time-consuming. Substances deemed promising based on results from the protein tests are then tested in lower throughput cellular and animal tests.

There have been some attempts to use image acquisition techniques to screen a large number of substances based upon biological cell information. One such attempt is described in International Application No. WO 98/38490 in the names of Dunlay, et al. Dunlay et al. generally describes a conventional image acquisition system. This conventional system collects and saves images based on certain criteria that are predefined, not on a fixed area of an imaging surface. Additionally, the conventional system has poor lighting design, which makes image processing for multiple cells difficult. Furthermore, the conventional system is not designed for capturing, populating and utilizing a large database design. The conventional system is designed for customized cellular assays, not as a tool for generation of a cellular informatics database. Without such database capabilities the conventional system cannot be used for screening, analyzing, and comparing large quantities of cells from multiple experiments on multiple days in a predictive, efficient and cost effective manner.

What is needed is a rapid assay to assess the activity of compounds against multiple drug targets simultaneously in a cellular context. What is also needed are techniques for finding the effects of substances on cell function based upon searching and analyzing cellular information.

SUMMARY OF THE INVENTION

According to at least one embodiment of the present invention, techniques for determining information about effects of potential substances on cells are provided. In another exemplary embodiment, the present invention provides a novel system including hardware, computer codes, user interfaces, and a database for acquiring, storing and retrieving cellular and substance information. The cells can include living, dead, or fixed cells or fractions of cells. The present invention enables, *inter alia*, researchers and/or scientists to identify promising candidates in the search for new and better medicines or treatments using, for example, a cellular informatics database.

According to the present invention, a computer program for identification and verification of biological properties of substances can include code that causes a sample of a substance to be administered to a cell. The code determines one or more features for two or more cell components, or markers, in the presence of the substance. The code can form one or more descriptors from the features. Descriptors can be formed by combining features of two or more cell components as identified using the markers. The code can then search one or more descriptors obtained from prior administered substances upon cells in order to locate descriptors having a relationship to the descriptors noted for the substance under study. The code predicts properties of the administered substance based upon the properties of the prior administered substances using the relationship between the descriptors. The code can provide for identifying properties of substances based upon effects on cell characteristics. Candidate drug mechanisms of action, potency, specificity, pharmacodynamic, and pharmacokinetic parameters, toxicity, and the like can be used as substance properties.

In a specific embodiment, the present invention provides a system for acquiring knowledge from cellular information. The system has a database comprising a database management module ("DBMS"). The system also has a variety of other modules, including a population module that is coupled to the DBMS and serves to categorize and store a plurality of features (including but not limited to cell size, distance between cells, cell population, as well as sub-cellular features such as organelle location, protein location and sub-cellular constituent location and

movement) from an image acquisition device into the database. The system has a translation module coupled to the DBMS for defining a descriptor from a set of selected features from the plurality of features. In a specific embodiment, the descriptor is for a known or unknown compound, e.g., drug. A prediction module is
5 coupled to the DBMS for selecting one of a plurality of a descriptors from known and unknown compounds from the database based upon a selected descriptor from a selected compound. The selected compound may be one that is useful for treatment of human beings or the like.

In a specific embodiment, the present invention provides a system for
10 populating a database with cellular information. The system includes a cell holder (e.g., multi-well plate, chip, microfluidic assembly, or other cell chamber) comprising a plurality of sites in a spatial orientation. Each of the sites is capable of holding a plurality of cells to be imaged. Note – the light guide is one embodiment, but we don't want to be limited to it.

15 According to one embodiment, the present system also has an illumination apparatus including a liquid light guide operably coupled to the imaging device for highlighting the plurality of cells in a relatively even spatial manner for image capturing and measurement purposes. Still further, the liquid light guide allows sub-elements (e.g., filter, lamp) of the illumination apparatus to be placed at a
20 remote location to prevent mechanical interference of the cell holder during image capturing. Alternative lighting methodologies may, with equal facility, be implemented.

The system also has an image-capturing device (e.g., charge coupled device camera, translation stage, shutter, microscope, software, shutter control) coupled to a
25 computing device (e.g., computer, network computer, work station, analog computing device, on-board image-processor, and laptop). The image-capturing device is adapted to capture at least one image in at least one of the plurality of sites. One some embodiments, multiple images can be captured, where each image represents a different cell component (or portion). The image-capturing device can be adapted to
30 convert the image into a digital representation, which highlights the feature or features of the one site.

A database storage device (e.g., relational database, object oriented database, mixed object oriented database) includes a database management element. The

database is coupled to the image capturing device. In a specific embodiment, the present system includes modules for feature extraction, generation of descriptions, and data preparation and analysis.

In a specific embodiment, the present invention provides a novel
5 system for determining an effect of a manipulation of a cell using one or more image frames. The system has a plate comprising a plurality of sites in a spatial orientation. Each of the sites is capable of holding a plurality of cells to be imaged. The system also has an image capturing device to capture a plurality of images of at least one site from the plurality of sites. The image capturing device is coupled to the computing
10 device. The system also has an image processing device to combine the plurality of images of at least one site or plurality of sites. The image processing device is operably coupled to the plate. An image processing device is also included. The image processing device can be adapted to form a digitized representation of the plurality of images from the site or plurality of sites. Furthermore, the system has a
15 database storage device comprising a database management element. The database can be adapted to retrieve the descriptor or descriptors of the plurality of features from the computing processing device and storing them in a selected manner.

In a specific embodiment, the present invention provides a system for capturing cellular information. The system also has an image acquisition system
20 comprising a charged coupled device camera adapted to capture an image of a plurality of manipulated cells in various stages of the cell cycle. The stages of the cell cycle are currently understood to include interphase, G0 phase, G1 phase, S phase, G2 phase, M phase, prophase, prometaphase, metaphase, anaphase, and telophase. The principles of the present invention specifically contemplate the application thereof on
25 additional cell cycle stages when and if they are identified.

An optical source is coupled to the image acquisition system for highlighting the plurality of manipulated cells in the various stages of the cell cycle. The illumination apparatus provides for an acquisition of the image of the plurality of manipulated cells. In a specific embodiment, the illumination apparatus has a liquid
30 light guide coupled to a light source at a remote location.

A variety of user interfaces are utile for accessing the several features of the present invention. Those having ordinary skill in the art will appreciate that different user interfaces may be required to support different research scenarios. The

present invention specifically contemplates the utilization of a wide variety of user interfaces.

Numerous benefits are achieved by way of the present invention over conventional techniques. The present invention can provide techniques for predictive cellular bioinformatics that can streamline a number of important decisions made in the drug discovery industry. The present invention can be implemented using off the shelf hardware including databases. In other aspects, the present invention can find useful information about substances as well as cells or portions of cells. Furthermore, the present invention can acquire more than one feature using more than one manipulation. Moreover, the present invention can provide information about a wide variety of cellular information that is not conventionally available. This information includes information about different cell components, e.g., nuclei and Golgi apparatus. Still further, the present invention provides an automated or semi-automated technique for acquiring images and populating a database. The present database can be combined with others such as genomics, and the like. Moreover, the present invention can be implemented to predict, *inter alia*, a mechanism of action, toxicity, target validation, and pre-clinical disease model.

A further understanding of the nature and advantages of the invention herein may be realized by reference to the remaining sections of the specification and the attached drawings.

BRIEF DESCRIPTION OF THE DRAWING

For more complete understanding of the present invention, reference is
5 made to the accompanying Drawing in the following Detailed Description of the
Invention. In the drawing:

Fig. 1 is a simplified system diagram according to an embodiment
according to the present invention;

10 Figs. 1A-1B are more detailed diagrams of database systems according
to embodiments of the present invention;

Fig. 2 is a simplified block diagram according to an alternative
embodiment according to the present invention;

Figs. 3-6 are simplified diagrams of system elements according to
embodiments of the present invention;

15 Figs. 7A-7K illustrate representative block diagrams of simplified
process steps in a particular embodiment according to the present invention;

Fig. 8A-8F illustrate representative quantified descriptors of effects of
manipulations on images of cells in a particular experiment;

20 Fig. 9 illustrates example images for different types of morphologies in
a particular experiment;

Fig. 10 illustrates a distribution of various morphologies in a cell
population responsive to drug concentration in a particular experiment;

Fig. 11 illustrates a graph of quantified features of effects of
manipulations on cells in a particular experiment;

25 Fig. 12 illustrates effects of external agents on cells in a particular
experiment;

Fig. 13 illustrates 4 panels for each marker for a plurality of A549 cells
in a particular experiment;

30 Fig. 14 illustrates 4 panels for each marker for a plurality of OVCAR-3
cells in a particular experiment;

Fig. 15 illustrates 4 panels for each marker for a plurality of OVCAR-3
cells at 20x in a particular experiment;

Fig. 16 illustrates 4 panels for each marker for a plurality of OVCAR-3 cells at 40x in a particular experiment;

Fig. 17 illustrates a representative input for a morphometric analysis program in a particular embodiment according to the present invention; and

5 Figs. 18-19 illustrate examples of the generation of pseudo-sequences and clustering in a particular embodiment according to the present invention.

Fig. 20 is a block diagram for a first research scenario;

Fig. 21 is a block diagram for a second research scenario; and

Fig. 22 is a block diagram for a third research scenario.

10 Reference numbers refer to the same or equivalent parts of the invention throughout the several figures of the Drawing.

DETAILED DESCRIPTION OF THE INVENTION

According to the present invention, techniques for determining information about manipulated cells or substances based upon living, fixed, or dead cell structures or portions of cells are provided. In an exemplary embodiment, the present invention provides a novel system including computer codes coupled to a database and user interfaces for acquiring, storing and retrieving such information. Other embodiments provide a novel image capturing system for providing digitized representations of live and dead cell structures or the like.

Fig. 1 is a simplified system diagram 10 of a cellular knowledge-based system according to an embodiment of the present invention. This diagram is merely an example and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. The present system 10 includes a variety of elements such as a computing device 13, which is coupled to an image processor 15 and is coupled to a database 21. The image processor receives information from an image capturing device 17, which image processor and image capturing device are collectively referred to as the imaging system herein. The image capturing device obtains information from a plate 19, which includes a plurality of sites for cells. These cells can be biological cells that are living, fixed, dead, cell fractions, cells in a tissue, and the like. The computing device retrieves the information, which has been digitized, from the image processing device and stores such information into the database. A user interface device 11, which can be a personal computer, a work station, a network computer, a personal digital assistant, or the like, is coupled to the computing device.

Fig. 1A is a simplified diagram of a database system 1000 according to an embodiment of the present invention. This diagram is merely an example and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize many other variations, modifications, and alternatives. Database system 1000 includes a variety of techniques for processing images from biological cells, e.g., fixed, living, and dead cells, and cell portions. As shown, images are acquired 1001. These images can be from a single frame or multiple frames. As merely an example, an image processing system may analyze such images. One example of

such an image processing system is described below, but should not be construed as limiting certain claims.

In a specific embodiment, cell samples are manipulated using a compound (e.g., substance, drug). The cell samples are imaged for a simple portion or portions, e.g., manipulated cell substructure, manipulated spatial feature of cell, cell density. Image processing techniques are used to extract 1003 the feature or features from the image or images. The features can be an independent or a dependent set of cell characteristics (which may be predominately visual) including, for example, count, area, perimeter, length, breadth, fiber length, fiber breadth, shape factor, elliptical form factor, inner radius, outer radius, mean radius, equivalent radius, 10 equivalent sphere volume, equivalent prolate volume, equivalent oblate volume, equivalent sphere surface, average intensity, total intensity, optical density, radial dispersion, texture difference, and others. Each of these features corresponds to a similar manipulation by a compound. Each manipulation forms a new set of features, which are identifiable to the compound. Once each set of features has been extracted, 15 the feature set is populated 1004 into a database 1012. Accordingly, the database includes many sets of features, where each set corresponds to a different manipulation for a selected cell. Each set of features corresponding to a manipulation provides a descriptor 1009, which is also stored 1019 in the database. The descriptor is a "finger print" including each feature for the manipulation. Each descriptor may be unique, or 20 may have similarities to other descriptors or may even be the same as other descriptors for known and unknown manipulations.

The present system retrieves features, which we define as simple features herein, and forms composite features 1007 from them. More than one feature 25 can be combined in a variety of different ways to form these composite features. In particular, the composite feature can be any function or combination of a simple feature and other composite features. The function can be algebraic, logical, sinusoidal, logarithmic, linear, hyperbolic, statistical, and the like. Alternatively, more than one simple feature can be combined in a functional manner (e.g., 30 arithmetic, algebraic). As merely an example, the composite feature equals a sum of feature 1 and feature 2, where these features correspond to the same manipulation. Alternatively, the composite feature equals feature 1 divided by feature 2. Alternatively, the composite feature equals feature 1 minus feature 2. Alternatively,

the composite feature equals a constant times feature 1 plus feature 2. Of course, there are many ways that the composite feature can be defined. The present system also stores 1017 these features in the database. The composite features can also be further combined with simple features. Once these features are defined as descriptors, they are stored 1019 in the database.

Fig. 1B is a simplified diagram of a database system engine 2000 according to an embodiment of the present invention. This diagram is merely an example and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize many other variations, modifications, and alternatives. The engine can be implemented into the present database for populating, searching, and predicting compound or cell characteristics. As merely an example, engine 2001 includes an input/output module 2008. The input/output module is used to input and output information from the database. The information includes, among others, a plurality of feature sets, which correspond to many manipulations. Additionally, the information includes descriptors, which each corresponds to a set of features from the manipulation. The database also has a population module, which is used to configure the features based upon an entity relationship, which has been predetermined.

The database engine also has other modules. In particular, the database has a transcription module, which transfers a preselected set of features and creates a descriptor from them. The transcription module can be used to take a known compound, which has features, to transcribe them into a descriptor. Alternatively, the transcription module can be used to take an unknown compound, which has features, to transcribe them into a descriptor. These descriptors are provided into the database for subsequent use. Finally, the database engine has a prediction module, which can be used to potentially predict a property (e.g., mechanism of action) of an unknown compound. Here, the unknown compound is provided with a descriptor, but the property of the compound is unknown. In one embodiment, the prediction module compares a descriptor of an unknown compound with the many descriptors of known compounds, which were in the populated database. Depending upon the matching criteria, the prediction module will attempt to uncover one or more descriptors of known compounds. Once the prediction module finds the descriptors of the known compounds based upon the descriptor for the unknown compound, it identifies a potential property of such unknown compound for analysis and review. Here, it is

believed that certain features of the known compound, which are similar to those features of the unknown compound may uncover a property to the unknown compound. Details of the present software engine are described more fully below.

Fig. 2 is a simplified block diagram 20 of a cellular knowledge-based system according to an alternative embodiment of the present invention. This diagram is merely an example and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. Like reference numerals are used in the present diagram as the previous diagram for easy cross-referencing, but are not intended to be limiting in any manner.

10 The present diagram 20 includes a variety of elements such as a processor 13 or computing device coupled to a database 11. The processor can be used for retrieving and storing information from the database. The system also includes a plurality of system elements, such as a cleaner 23, a dispenser 25, and an image capturing system 27, which are also coupled to the database in some embodiments. These elements can

15 be coupled to each other through a network or the like. As merely an example, the network can be a NetWareTM network from Novell Corporation or an internet network or the Internet but can also be others and any combination thereof. The system also has an output device 31, which can be used to output information from the database, processor, or other system elements. Details of these elements are described more

20 fully below in reference to the Figs.

Figs. 3-5 are simplified drawings of system elements according to embodiments of the present invention. These diagrams are merely examples and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. As merely an example,

25 Fig. 3 is a simplified diagram of a processor or computing device 13. The computing device 13 includes a bus 112 which interconnects major subsystems such as a central processor 114, a system memory 116 (e.g., random access memory), an input/output ("I/O") controller 118, an external device such as a display screen 124 via a display adapter 126, a keyboard 132 and a mouse 146 via an I/O controller 118, a SCSI host

30 adapter (not shown), and a floppy disk drive 136 operative to receive a floppy disk 138.

The computing device has other features. Storage Interface 134 may act as a storage interface to a fixed disk drive 144 or a CD-ROM player 140 operative

to receive a CD-ROM 142. Fixed disk 144 may be a part of computing device or may be separate and accessed through other interface systems. A network interface 148 may provide a direct connection to a remote server via a telephone link or to the Internet. Network interface 148 may also connect to a local area network ("LAN") or other network interconnecting many computer systems. Many other devices or subsystems (not shown) may be connected in a similar manner. Also, it is not necessary for all of the devices shown in Fig. 3 to be present to practice the present invention, as discussed below. The devices and subsystems may be interconnected in different ways from that shown in Fig. 3. The operation of a computer system such as that shown in Fig. 3 is readily known in the art and is not discussed in detail in this application. Computer code to implement the present invention, may be operably disposed or stored in computer-readable storage media such as system memory 116, fixed disk 144, CD-ROM 140, or floppy disk 138. The computer code can be organized in terms of processes or modules, depending upon the application. That is, the computer code can include a prediction module, a translation module, or other modules to carry out the functionality described herein, as well as others.

Figs. 4 and 5 are simplified diagrams of an imaging system 200 according to an embodiment of the present invention. As shown, the imaging system 200 includes a variety of features such as housing 203, which holds a stage assembly 204. The stage assembly includes an x-stage movement element 206, which is along an x-direction, and a y-stage movement element 207, which is along a y-direction. The imaging system also includes a z-direction movement element, which is perpendicular to the x-y plane. The z-direction movement motor can be attached to the stage, or to the objective nosepiece by way of the microscope housing, or as an external motor between the objective and the microscope housing. The stage can align in any one of the directions to an accuracy of one micron and less, or one-half micron and less, or one-quarter micron and less, depending upon the embodiment.

The stage holds a plate 202 or cell holder, which houses one of a plurality of samples. The plate includes a spatial array 209 of process sites. Each of the process sites can include a plurality of cells and solutions depending upon the embodiment. Each of the sites can carry a sufficient amount of solution to prevent substantial evaporation of the sample during processing in some embodiments. In embodiments for large scale analysis, the plate includes at least 96 sites, or more than

or equal to 384 sites, or more than or equal to 1,536 sites. The plate bottom is transparent and thin, which allows light to pass through the sample. Additionally, the plate is made of a suitable chemical resistant material. As merely an example, the plate can be either a 96, or 384, or 1536 or other formats from places such as Becton Dickinson of Franklin Lakes, NJ, or Corning Science Products of Corning, NY. In a preferred embodiment, the plate is a Corning Costar black-walled 96 well plate catalog #3904 from Corning Science Products of Corning, NY, but should not be limited to these in some applications, but can be others.

Also shown is the condenser for the microscope 201, which can be used to collect phase, DIC, or bright field images of the cells. Images resulting from the illumination of the samples to fluorescence, phase, DIC, or bright field techniques are collected using an image capturing device 208, which captures an image or images of cells from the plate. In a specific embodiment, the microscope is an inverted configuration with the objectives on the bottom of the plate and the condenser disposed overlying an upper surface of the sites, while the image capturing device underlies the sites. Images captured by the imaging device, whether analogue or digital, are viewed by a monitor or other devices. The image capturing device can be any camera assembly such as a charge coupled device camera, which is known as a CCD camera, or other high resolution camera capable of capturing images from the sites. In a specific embodiment, the camera is an interline CCD camera which does not require an external shutter.

In a specific embodiment, the present imaging system can be any suitable unit that is flexible for automated image collection using multi-well plastic plates. The imaging system also should be adapted to collect high-resolution images of cells on plastic or glass plates, cell growth chambers, or coverslips. The system also can be used for imaging multiple cell markers in multiple imaging conditions. To accomplish this, the microscope system has a variety of elements such as a light source, a motorized excitation filter wheel and shutter, x-y-z-motorized stage, excitation and emission filters, Fluor phase and DIC objectives, motorized objective nosepiece, dichroic filters, motorized dichroic filter cubes, phase and DIC rings and prisms, CCD camera, and software control. As merely an example, the present imaging system can have components such as those listed in the Table below.

DESCRIPTION	MAKER	MODEL
Microscope	Zeiss	100M
(x-y) motorized stage	Prior	
Xenon lamp	Sutter	Lambda
Filter wheel	Sutter	Lambda-10
Microtitre Plate holder	Prior	500-H223R
Isolation Table	Kinetic Systems	9101-24-85
Objective Spacers	Polytec PI	P-721.90
Camera	Hamamatsu	C47-95
Computer	IBM	IntelliStation
Software	Metamorph	v.4
Objectives	Zeiss	Achroplan 10x/0.25 LD-Achroplan 20x/0.4 LD-Achroplan 40x/0.6

Table: Image Acquisition System Elements

5 In a specific embodiment, the present system has the following capabilities, which are not intended to be limiting.

Image acquisition

- 1) Ability to automatically acquire multi-wavelength images from multiple sites on one multi-well plate, to sequentially name image files, and to log any
10 imaging parameter information with image files.
- 2) Ability to link images with a larger database/spreadsheet of information.
- 3) Ability to automatically collect multiple plates by interfacing the imaging system with a robotic arm.

15

X-Y control

- 1) Ability to place 96, 384, or 1536 well plates onto microscope stage and move to each well sequentially.

2) Ability to return to each well and collect another round of images (multi-site time-lapse) or ability to collect rapid time-lapse information at each well (time-lapse of many wells).

3) Ability to collect a low magnification image, automatically determine features which may be of interest, automatically change the objective to a higher magnification, and collect high magnification images of a fixed number of those identified cells in the sample.

4) Ability to collect multiple frames in each site.

10 Z control

1. Ability to auto-focus with substantially minimal damage to biological specimen or fluorophore.

2. Ability to auto-focus rapidly.

15 The present embodiment of the imaging system is shown by way of Figs. 5A and 5B. These diagrams are merely examples and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. The present imaging system 40 includes a variety of elements such as a microscope 41, which is preferably an epi-fluorescent microscope, but can be confocal, multiphoton, or hybrid types. The microscope includes elements
20 41A, the motorized Z-axis; 41B, the motorized dichroic filter cube holder; and 41C, the motorized objective nosepiece. In one embodiment, the microscope is a Model 100M made by Zeiss. The microscope communicates to computer 51 through control lines 73, 75, and 76. The imaging system also has camera 50 coupled to controller
25 50A and computing device 51, which oversees and controls operations of the elements of the imaging system.

The present microscope includes drivers for spatially moving a stage in two dimensions, including an x-direction, a y-direction, and moving the objective nosepiece in a z-direction in a Cartesian coordinate system. The z-direction
30 movement is provided using a fast z-motor, which can make z-direction adjustments within a predetermined time. The z-direction movement generally provides for focussing of the sample to the camera. The focussing occurs within the predetermined time of preferably ten seconds and less, or five seconds and less, or one

second and less, depending upon the embodiment. As merely an example, the z-motor or positioner can be a model PIFOC objective nanopositioner made by a company called Physik Instrumente of Waldbronn, Germany, but also can be others. The z-motor couples to computer 51 through line 63, which may also include a
5 controller. Depending upon the embodiment, a second z-motor 41A connected to the computer 51 by line 73 may be used to keep the z-motor 42 in the center of its travel. Alternatively, in other embodiments the stage could be provided with a z-motor allowing for movement of the stage in the z-direction.

The present stage also includes an x-y stage 43. The x-y stage moves
10 plate 59, e.g., 96 site, 384 site, 1536 site. The x-y stage moves plate in an x-y spatial manner. The stage has an accuracy or repeatability of about 1 micron and less, or about 2 microns and less. The stage can move in a continuous manner or a stepped manner. The stage also can move up to 30 mm/sec. or faster. The stage also can move 1 mm/sec. and less, depending upon the embodiment. The stage can also step
15 0.1 micron and less or 1 micron and less, as well as other spatial dimensions. The stage can be one such as a Proscan Series made by Prior Scientific of Rockland, MA but can also be others. The stage is controlled via control line 61 through controller 43A, which couples to computer 51 through control line 65.

The stage includes plate holder 44. The plate holder can hold a single
20 plate. In other embodiments, plate holder can also hold multiple plates. The plate holder can use mechanical, electrical, fluid, vacuum and other means for holding the plate or plates. The plate holder also is sufficiently stable for securing the plate. As merely an example, the plate holder is a Model 500-H223R made by Prior Scientific of Rockland, MA. In some embodiments, the plate holder may need adjustment in
25 the z-direction to provide for a desirable focus of a sample on a plate. In these embodiments, the plate holder is supported by spacers 45 or a plurality of stage pins, which mechanically elevate the plate holder in the z-direction. These pins are generally made of a suitable material for supporting such plate holder and also are sufficiently resistant to chemicals and the like.

30 In some embodiments, the entire imaging system is placed on an isolation table 49. The isolation table is disposed between the microscope and support structure. The isolation table is designed to prevent excessive vibration of the plate. The isolation table is made of a suitable material such as steel and honeycomb but can

be others. The table has a thickness of about 8 inches or preferably less than about 24 inches. In one embodiment, the table is Model 9101-24-85 made by Kinetic Systems of Boston, MA.

The imaging system also has a lamp or illumination assembly 62. The
5 lamp assembly provides for a light source (See reference letter B) to a plurality of elements in the imaging system. For easy reading, the light path is defined by the dotted lines, which are not intended to be limiting. The lamp assembly has a variety of elements such as a Xenon lamp 46. The Xenon lamp provides light at about 320 to 700 nanometers (Prefocused). The Xenon lamp is 175 or 300 Watts. As merely an
10 example, the lamp can be a Lambda Model made by Sutter Instrument Company of Novato, CA.

Referring to Fig. 5B, the lamp assembly also has a cold mirror 58, an excitation filter wheel 48, excitation filter(s) 55, and an excitation light shutter 57. As shown, light is derived from the Xenon lamp, reflects off of the cold mirror 58,
15 traverses through the excitation filter or filters 55, and is controlled by the excitation light shutter 57. The lamp assembly has filter wheel 48, which houses one of a plurality of filters, including excitation filters. The shutter and filter wheel are controlled via control lines 67, which are coupled to a computer 51 or other type of computing device. The control lines 67 are coupled through controller 57A (for
20 element 57) and controller 48A (for element 48) via control line 69 to computer 51.

Preferably, light traverses from the lamp assembly through a light guide 47 to illuminate features within the plate. The light guide is suitably selected to have a flexible member, which can be used to place lamp source at a remote location away from the imaging device. The flexible member substantially keeps any
25 vibration from the lamp assembly away from the imaging device. In some embodiments, the member is at least 1 foot away from the imaging device. The light guide is a guide, which is a flexible hose-type sleeve. The sleeve is filled with a liquid such as an aqueous solution containing chloride or phosphate. A thin layer may be formed on the inside of the sleeve. The layer can be a containing
30 tetrafluoroethylene and hexafluoropropylene, or containing tetrafluoroethylene and perfluoromethyl vinyl ether, or tetrafluoroethylene and perfluoropropyl vinyl ether. An example of such a light guide is described in International Application No. WO/98/38537 filed February 29, 1997, and assigned to NATH, Gunther. The liquid

light guide has less than about 30% transmission loss of the light at a remote location such as the imaging system.

Light is derived from the lamp assembly and directs off of filter 56, which directs the light upward. Filter 56 can be a dichroic and emission filter, as well as others. The light traverses through microscope nosepiece 41C, and traverses
5 through objective spacers 54. An objective 53 magnifies the light toward a predetermined point on the plate 59. The objective can be, for example, made by Zeiss of Jena, Germany, as well as other companies. The objective can be one of a plurality including 1X, 10X, 20X, 40X, and others, depending upon the application.
10 Magnification can be further expanded or contracted by intermediate optics between the objective and the camera. Selection of filter or filters is controlled by computer 51 via control line 75.

The camera 50 captures an image of cells from plate 59. The image is obtained from light scattering off of cells or portions of cells in the plate through
15 objective 53, through objective spacers, through filters 56, which are captured at camera 50. In this preferred embodiment, the camera is a digital camera, but can be an analogue camera. The digital camera is a CCD camera, which has 1280 by 1024 pixels, or more or less. The pixels can be 6.7 microns in dimension or more or less. The camera preferably is substantially free from an external shutter to quickly capture
20 a plurality of images of cells from the plate. The camera is controlled via control line 71 through controller 50A, which connects to computer 51 through control line 70. The present invention can also include other types of image acquisition devices selected from at least an epifluorescence, a confocal, a total-internal reflection, a phase, a Hoffman, a bright field, a dark field, a differential interference contrast, an
25 interference reflection, or multi-photon illumination device.

The present imaging system stores images on a high density memory device 60. The high density memory device is preferably optical, but can also be magnetic. The high density memory device can be any suitable unit that is capable of storing a plurality of images from a plurality of sites in the plate. The memory device
30 can be a compact disk, which would generally use a compact disk burner or the like. Depending upon the embodiment, the high density memory device is used to archive the images that are captured from the camera in the imaging system. Further details

of the imaging system can be found throughout the present specification, and more particularly below.

As merely an example, the present invention can be implemented using the following sequence of steps, which have been described in a journal entry form.

- 5 Here, images are opened and objects are identified based on a background value that has been edited in starting image acquisition. Information is maintained in a spreadsheet or other database format, which has the following information for each object:

Image Name	Image Plane	Image Date and Time
Elapsed Time	Object #	Total area
Pixel area	Area	Hole area
Relative hole area	Standard area count	Perimeter
Length	Breadth	Fiber length
Fiber breadth	Shape factor	Ell. form factor
Inner radius	Outer radius	Mean radius
Average gray value	Total gray value	Optical density
Radial dispersion	Texture Difference Moment	EFA Harmonic 2, Semi-Major Axis
EFA Harmonic 2, Semi-Minor Axis	EFA Harmonic 2, Semi-Major Axis Angle	EFA Harmonic 2, Ellipse Area
EFA Harmonic 2, Axial Ratio	EFA Harmonic 3, Semi-Minor Axis	

10

After computations are done, the log file is saved. In particular, the file is saved in an appropriate place with an appropriate name.

In a specific embodiment, the present invention provides the following detailed example of journal entries, which should not limit the scope of the invention.

Set Up Sequential File Names	Interactive: user sets up prefix name and image storage directory
Open Data Log	Opens a DDE (Excel) File
Annotate Log File	Interactive: experimental information that will go into the first line of the log file of stage positions
Stage (Go to Origin)	Origin is set as the center of well A1
Stage (Move to Absolute Position)	Offset to upper left hand corner of well (1410, 1621)
Stage (Log Position)	
Stage (Scan Wells)	User picks wells to scan: runs 3x3 image collection.jnl.

3X3 IMAGE COLLECTION.jnl

Stage (Scan)	Takes 9 images of well, -1600 motor steps apart from left to right 3 columns and 3 rows, runs FOCUS, COLLECT IMAGE, SAVE SEQUENTIAL FILE NAME.JNL.
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5

FOCUS, COLLECT IMAGE, SAVE SEQUENTIAL FILE NAME.jnl.

Stage (Log Position)	Logs stage position of each image
ADC – Focus	Opens up the manual focusing window with whatever focus time is current set
Show Message and Wait	Interactive: user hits enter to continue when done focusing

ADC-Acquire from Digital Camera	Takes Hoechst image
Save Using Sequential File Names	
Close	Closes image window

START IMAGE ANALYSIS.jnl

Low Pass	3x3 convolution of already opened image
Low Pass	3x3
Show Region Statistics	Interactive: Show entire image statistics. Calculate background subtraction value for step 4. by: INTENSITY Average + INTENSITY Std. Dev.
Arithmetic	Interactive: User inputs subtraction value from 3. into the constant Value field
Threshold image	Creates threshold 1 unit above 0 to 4096
Integrated Morphometry – Load State	Loads Start Image Analysis.ima Classifier 100 < area < 200000
Integrated Morphometry – Measure	Interactive: Shows area summary information about all objects. The average number is used as the Standard Area in 8.
Object Standards - Set Object Standards	Interactive: User inputs average area value from 7. into Standard Area box to be used by automated IMA for all images

IMA OBJECTS.jnl

Low Pass	3x3 convolution
Low Pass	3x3 convolution
Arithmetic	This background subtraction value needs to be manually entered into this journal from the value determined in START IMAGE ANALYSIS.jnl step 3
Threshold Image	1 unit above 0
Integrated Morphometry – Load State	Hoechst.IMA Classifier $200 < \text{area} < 200000$
Integrated Morphometry – Measure	Measures statistical info for all objects
Run Journal	Runs log obj and sum data.jnl

Log obj and sum data.jnl

Integrated Morphometry – Log Data	Logs object data into Sheet 1
Integrated Morphometry – Log Data	Log summary data into Sheet 2

5

COLLECT AUTOMATED IMA DATA IN ONE SPREADSHEET.jnl

Run Journal	Runs OPEN OBJECT LOG DDE FILE.JNL
Loop for all Images in a Directory	Loops IMA OBJECTS.jnl
Close Summary Log	
Close Object Log	User must manually save Excel spreadsheet

OPEN OBJECT LOG DDE FILE.jnl

Open Object Log	Opens a DDE object log into sheet 1 of an Excel spreadsheet
Open Summary Log	Opens a summary log into sheet 2

COLLECT AUTOMATED IMA DATA IN ONE SPREADSHEET 16 BIT IMAGES.jnl

Arithmetic	Interactive: Opens Arithmetic window for user to input background subtraction level from START IMAGE ANALYSIS.jnl step 3
Run Journal	Runs OPEN OBJECT LOG DDE FILE.JNL
Loop for all Images in a Directory	Interactive: Runs IMA OBJECTS 16 bit.jnl. User picks directory from which to choose.

5

IMA OBJECTS 16bit.jnl

Low Pass	3x3 convolution
Low Pass	3x3 convolution
Copy to 8-bit Image	No autoscale, to new untitled image
Save Using Sequential File Name	Saves 8bit image using previously defined Sequential File names.
Arithmetic	This background subtraction value needs to be manually entered into this journal from the value determined in START IMAGE ANALYSIS 16 TO 8 BIT.jnl step 5
Threshold Image	1 unit above 0 to 255

Integrated Morphometry – Load State	Hoecsht.IMA Classifier 200 < area < 200000
Integrated Morphometry – Measure	Measures statistical info for all objects
Run Journal	Runs log obj and sum data.jnl

START IMAGE ANALYSIS 16 to 8 BIT.jnl

Copy to 8-bit Image	No autoscale, to new untitled image
Close	Closes 16 bit image
Low Pass	3x3 convolution
Low Pass	3x3 convolution
Show Region Statistics	Interactive: Show entire image statistics. Calculate background subtraction value for step 6. by: INTENSITY Average + INTENSITY Std. Dev.
Arithmetic	Interactive: User inputs subtraction value from 5. into the constant Value field
Threshold image	Creates threshold by 1 unit above 0 to 255
Integrated Morphometry – Load State	Loads Start Image Analysis.ima Classifier 100 < area < 200000
Integrated Morphometry – Measure	Interactive: Shows area summary information about all objects. The average number is used as the Standard Area in 10.
Object Standards - Set Object Standards	Interactive: User inputs average area value from 9. into Standard Area box to be used by automated IMA for all images

IMA OBJECTS WITH NEW LOG FILE.jnl

Run Journal	OPEN OBJECT LOG DDE FILE.JNL
Run Journal	IMA OBJECTS.jnl
Close Summary Log	
Close Object Log	User must manually save every Excel spreadsheet generated.

INTERACTIVE IMA OBJECTS.jnl

Threshold Image	User manually sets threshold
Integrated Morphometry – Load State	Hoechst.IMA Classifier 200 < area < 200000
Integrated Morphometry – Measure	Objects
Integrated Morphometry – Log Data	Into open object.log file

5

COLLECT INTERACTIVE IMA DATA.jnl

Close Object Lo g	
Open Object Log	Interactive
Annotate Log File	Interactive: experimental information that will go into the first line of the object log file
Loop for all Images in Directory	Runs INTERACTIVE IMA OBJECTS.jnl

CHANGE FILTER, COLLECT IMAGE. SAVE SEQUENTIAL FILE
NAME.jnl

Stage (Log Position)	
ADC-Focus	

Show Message and Wait	Interactive – user presses Enter when done focusing
ADC – Acquire from Digital Camera	Hoechst
Save Using Sequential File Name	
Close	Close open image

COLLECT HOECHST AND FITC.jnl

Run Journal	FOCUS, COLLECT IMAGE, SAVE SEQUENTIAL FILE NAME.JNL
Run Journal	CHANGE FILTER, COLLECT IMAGE, SAVE SEQUENTIAL FILE NAME.jnl

3X3 IMAGE COLLECTION HOECHST FITC.jnl

Stage (Scan)	COLLECT HOECHST AND FITC.jnl
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5

AUTOMATED 3X3 IMAGE COLLECTION HOECHST FITC.jnl

Set Up Sequential File Names	Interactive: user sets up prefix name and image storage directory
Open Data Log	Excel DDL files
Annotate Log File	Interactive: experimental information that will go into the first line of the log file of stage positions
Stage (Go to Origin)	Origin is set as the center of well A1
Stage (Move to Absolute Position)	Offset to upper left hand corner of well (1410, 1621)

Stage (Log Position)	
Stage (Scan Wells)	Interactive: user picks wells to scan: runs 3X3 IMAGE COLLECTION HOECHST FITC.jnl

AUTOMATED IMAGE COLLECTION.jnl

Set Up Sequential File Names	Interactive: user sets up prefix name and image storage directory
Open Data Log	Opens a DDE (Excel) File
Annotate Log File	Interactive: experimental information that will go into the first line of the log file of stage positions
Stage (Go to Origin)	Origin is set as the center of well A1
Stage (Log Position)	
Stage (Scan Wells)	Interactive: user picks wells to scan: runs FOCUS, COLLECT IMAGE, SAVE SEQUENTIAL FILE NAME.JNL. Well to well travel = (-9035, -9035)

5

STARTUP.jnl

Install and Configure Devices	Open Stage Meta Devices
Set Live Video Channel	

Preferences	<u>Measure Objects</u> : Draw failed classifier objects, Exclude objects that touch the edge of the image, Enable Elliptical Fourier Parameters, turn off Warn users when measurement data will be erased <u>Image Saving</u> : Save Tiff/stk using LZW compression <u>Image Windows</u> : Use transparent thresholds.
Configure Default Paths	C:\Metamorph Data C:\Metamorph Data\Commmon Settings
Load Journal Taskbar	Common.JTB

Nested Journals

Automated 3x3 Image Collection

5 *Loop* 3x3 image collection
 Loop focus, collect image, save sequential file name

Automated 3x3 image collection Hoechst FITC

10 *Loop* 3x3 image collection Hoechst FITC
 loop Collect Hoechst and FITC
 focus, collect image, save sequential file name
 change filter, collect image, save sequential file name

Automated image collection

15 *Loop* focus, collect image, save sequential file name

Collect automated IMA data in one Spreadsheet

Open object log DDE file

Loop IMA objects

Log obj and sum data

Collect automated IMA data in one spreadsheet 16 bit images

5 Open object log DDE file

Loop IMA objects 16 bit

Log obj and sum data

10 Although the above has been generally described in terms of a specific user interface and software code, other user interfaces and code can also be used. One of ordinary skill in the art would recognize many other variations, alternatives, and modifications.

Fig. 6 is a simplified diagram 600 of a cleaning and dispensing system according to an embodiment of the present invention. This system 600 includes a
15 variety of elements such as a dispensing head 609, which is coupled to a plurality of pipettes 601. The pipettes input and output fluids or solutions from plate 603. The plate has a plurality of sites, each of which can be used to input cells or a combination of cells and solution. The system also has elements to house solutions 605, which are used to manipulate cell samples in the plate. The dispensing head is supported
20 through a support member 607, which is sufficiently rigid to allow for movement of the head. The dispenser is coupled to the present system in a mechanical and electrical manner, which provides for a fully integrated system for providing cell samples to the imaging system according to the present invention.

Fig. 7A illustrates a representative block flow diagram of simplified
25 process steps of a method for determining properties of a manipulation based upon effects of the manipulation on one or more portions of one or more cells in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. In step
30 700, one or more samples of cells can be provided. These cells can be live, dead, or fixed cells, or cell fractions. The cells also can be in one of many cell cycle stages, including G0, G1, S, G2 or M phase, M phase including the following cell cycle stages: interphase, prophase, prometaphase, metaphase, anaphase, and telophase.

Cell components tracked in presently preferable embodiments can include proteins, protein modifications, genetically manipulated proteins, exogenous proteins, enzymatic activities, nucleic acids, lipids, carbohydrates, organic and inorganic ion concentrations, sub-cellular structures, organelles, plasma membrane, adhesion complex, ion channels, ion pumps, integral membrane proteins, cell surface
 5 receptors, G-protein coupled receptors, tyrosine kinase receptors, nuclear membrane receptors, ECM binding complexes, endocytotic machinery, exocytotic machinery, lysosomes, peroxisomes, vacuoles, mitochondria, Golgi apparatus, cytoskeletal filament network, endoplasmic reticulum, nuclear membrane, proteosome apparatus,
 10 chromatin, nucleolus, cytoplasm, cytoplasmic signaling apparatus, microbe specializations and plant specializations.

The following table illustrates some markers and cell components commonly used by embodiments according to the present invention. Other markers can be used in various embodiments without departing from the scope of the
 15 invention.

Cell component	Marker	Disease State
Plasma membrane (including overall cell shape)	Carbocyanine dyes Phosphatidylserine Various lipids Glycoproteins	Apoptosis-Cancer Apoptosis-Neural degenerative Ds
Adhesion complexes	Cadherins Integrins Occludin Gap junction ERM proteins CAMs Catenins Desmosomes	Thrombosis Metastasis Wound healing Inflammatory Ds Dermatologic Ds
Ion Channels and Pumps	Na/K Atpase Calcium channels Serotonin reuptake pump CFTR	Cystic fibrosis Depression Congestive Heart Failure Epilepsy

G coupled receptors	β adrenergic receptor Angiotensin receptor	Hypertension Heart Failure Angina
Tyrosine kinase receptors	PDGF receptor FGF receptor IGF receptor	Cancer Wound healing Angiogenesis Cerebrovascular Ds
ECM binding complexes	Dystroglycan Syndecan	Muscular Dystrophy
Endocytotic machinery	Clathrin Adaptor proteins COPs Presenilins Dynamin	Alzheimer's Ds
Exocytotic machinery	SNAREs Vesicles	Epilepsy Tetanus Systemic Inflammation Allergic Reactions
Lysosomes	Acid phosphatase Transferrin	Viral diseases
Peroxisomes/Vacuoles		Neural degenerative Ds
Mitochondria	Caspases Apoptosis inducing factor F1 ATPase Fluorescein Cyclo-oxygenase	Apoptosis Neural degenerative Ds Mitochondrial Cytopathies Inflammatory Ds
Golgi Apparatus	Lens Culinaris DiOC6 carbocyanine dye COPs	

Cytoskeletal Filament Networks	Microtubules Actin Intermediate Filaments Kinesin, dynein, myosin Microtubule associated proteins Actin binding proteins Rac/Rho Keratins	Cancer Neural degenerative Ds Inflammatory Ds Cardiovascular Ds Skin Ds
Endoplasmic Reticulum	SNARE PDI Ribosomes	Neural degenerative Ds
Nuclear Membrane	Lamins Nuclear Pore Complex	Cancer
Proteosome Apparatus	Ubiquityl transferases	Cancer
Chromatin	DNA Histone proteins Histone deacetylases Telomerases	Cancer Aging
Nucleolus	Phase markers	
Cytoplasm	Intermediary Metabolic Enzymes BRCA1	Cancer
Cytoplasmic Signaling Apparatus	Calcium Camp PKC pH	Cardiovascular Ds Migraine Apoptosis Cancer
Microbe Specializations	Flagella Cilia Cell Wall components: Chitin synthase	Infectious Ds

Plant specializations	Choloroplast Cell Wall components	Crop Protection
-----------------------	--------------------------------------	-----------------

Then, in a step 702, one or more samples of the manipulation can be provided to the cells. Manipulations can comprise one or any combination of chemical, biological, mechanical, thermal, electromagnetic, gravitational, nuclear, or temporal factors, for example. For example, manipulations could include exposure to chemical compounds, including compounds of known biological activity such as therapeutics or drugs, or also compounds of unknown biological activity. Or exposure to biologics that may or may not be used as drugs such as hormones, growth factors, antibodies, or extracellular matrix components. Or exposure to biologics such as infective materials such as viruses that may be naturally occurring viruses or viruses engineered to express exogenous genes at various levels. Bioengineered viruses are one example of manipulations via gene transfer. Other means of gene transfer are well known in the art and include but are not limited to electroporation, calcium phosphate precipitation, and lipid-based transfection. Manipulations could also include delivery of antisense polynucleotides by similar means as gene transfection. Other genetic manipulations include gene knock-outs or gene mutations. Manipulations also could include cell fusion. Physical manipulations could include exposing cells to shear stress under different rates of fluid flow, exposure of cells to different temperatures, exposure of cells to vacuum or positive pressure, or exposure of cells to sonication. Manipulations could also include applying centrifugal force. Manipulations could also include changes in gravitational force, including sub-gravitation (the preferred embodiment in outer space). Manipulations could include application of a constant or pulsed electrical current. Manipulations could also include irradiation. Manipulations could also include photobleaching which in some embodiments may include prior addition of a substance that would specifically mark areas to be photobleached by subsequent light exposure. In addition, these types of manipulations may be varied as to time of exposure, or cells could be subjected to multiple manipulations in various combinations and orders of addition. Of course, the type of manipulation used depends upon the application.

Then, in a step 704, one or more descriptors of a state in the portions of the cells in the presence of the manipulation can be determined using the images

collected on the imaging system. Descriptors can comprise scalar or vector values, representing quantities such as area, perimeter, dimensions, intensity, gray level, aspect ratios, and the like. Other types of descriptors include, but are not limited to, one or any combination of characteristics such as a cell count, an area, a perimeter, a length, a breadth, a fiber length, a fiber breadth, a shape factor, a elliptical form factor, an inner radius, an outer radius, a mean radius, an equivalent radius, an equivalent sphere volume, an equivalent prolate volume, an equivalent oblate volume, an equivalent sphere surface area, an average intensity, a total intensity, and an optical density. These descriptors can be average or standard deviation values, or frequency statistics from the descriptors collected across a population of cells. These descriptors can be further reduced using other methods such as principal component analysis and the like. In some embodiments, the descriptors include features from different cell portions or cell types. That is, a first feature can be from a nuclei and a second feature is from another cell structure such as Golgi apparatus, mitochondria, spacing between cell structures or cells themselves, as well as many others.

A presently preferable embodiment uses descriptors selected from the following table. Other descriptors can also be used without departing from the scope of the invention.

Name of Parameter	Explanation/Comments
Count	Number of objects
Area	
Perimeter	
Length	X axis
Width	Y axis
Shape Factor	Measure of roundness of an object
Height	Z axis
Radius	
Distribution of Brightness	
Radius of Dispersion	Measure of how dispersed the marker is from its centroid
Centroid location	x-y position of center of mass
Number of holes in closed objects	Derivatives of this measurement might include, for

	example, Euler number (= number of objects - number of holes)
Elliptical Fourier Analysis (EFA)	Multiple frequencies that describe the shape of a closed object
Wavelet Analysis	As in EFA, but using wavelet transform
Interobject Orientation	Polar Coordinate analysis of relative location
Distribution Interobject Distances	Including statistical characteristics
Spectral Output	Measures the wavelength spectrum of the reporter dye. Includes FRET
Optical density	Absorbance of light
Phase density	Phase shifting of light
Reflection interference	Measure of the distance of the cell membrane from the surface of the substrate
1,2 and 3 dimensional Fourier Analysis	Spatial frequency analysis of non closed objects
1,2 and 3 dimensional Wavelet Analysis	Spatial frequency analysis of non closed objects
Eccentricity	The eccentricity of the ellipse that has the same second moments as the region. A measure of object elongation.
Long axis/Short Axis Length	Another measure of object elongation.
Convex perimeter	Perimeter of the smallest convex polygon surrounding an object
Convex area	Area of the smallest convex polygon surrounding an object
Solidity	Ratio of polygon bounding box area to object area.
Extent	proportion of pixels in the bounding box that are also in the region
Granularity	
Pattern matching	Significance of similarity to reference pattern
Volume measurements	As above, but adding a z axis

Then, in a step 705, a database of cell information can be provided. Next, in a step 706, a plurality of descriptors can be searched from a database of cell information in order to locate descriptors based upon one of the descriptors of the manipulation. Then, in a step 708, properties of the manipulation are predicted based
5 upon the properties of the located descriptors. Properties can comprise toxicity, specificity against a subset of tumors, mechanisms of chemical activity, mechanisms of biological activity, structure, adverse biological effects, biological pathways, clinical effects, cellular availability, pharmacological availability, pharmacodynamic properties, clinical uses and indications, pharmacological properties, such as
10 absorption, excretion, distribution, metabolism and the like.

In a particular embodiment, step 706 comprises determining matching descriptors in the database corresponding to a prior administration of the manipulation to the descriptors of the present administration of the manipulation. In a particular embodiment according to the present invention, combinations of
15 measurements of scalar values can provide predictive information. A database can be provided having one or more "cellular fingerprints" comprised of descriptors of cell-substance interactions of drugs having known mechanisms of action with cells. Such descriptors can be analyzed, classified, and compared using a plurality of techniques, such as statistical classification and clustering, heuristic classification techniques, a
20 technique of creating "phylogenetic trees" based on various distance measures between descriptors from various drugs. In this embodiment, numeric values for the descriptors can be used by comparison techniques. A phylogenetic tree can be created that illustrates a statistical significance of the similarity between descriptors for the drugs in the database. Because the drugs used to build the initial database are of
25 known mechanism, it can be determined whether a particular scalar value in a descriptor is statistically predictive. Finally, a compound descriptor with no known mechanism of action can be queried against the database and be statistically compared and classified among the drugs in the database that the compound most resembles.

In a particular embodiment, relationships between measured
30 morphological properties of images and physiological conditions can be determined. Relationships can include, for example, treatment of different cell lines with chemical compounds, or comparing cells from a patient with control cells, and the like. In a presently preferable embodiment, comparisons can be performed on acquired image

features. Some embodiments can comprise statistical and neural network - based approaches to perform comparisons of various features. The foregoing is provided as merely an example, and is not intended to limit the scope of the present invention. Other techniques can be included for different types of data.

5 In some embodiments, classification, clustering and other types of predictive data analysis can be performed on features extracted from cell images. In a presently preferable embodiment, statistical procedures for comparisons, classification and clustering are performed on data obtained from imaging cells.

Fragments of data preparation and pre-formatting (S language):

```
10       >tmp.frame <- Generic.Summary  
      >names1 <- paste("Cell.line.5", tmp.names, sep=".")  
      > by.compound.matrix <- as.matrix(arranged.by.compound)
```

 Example of the code for principal component analysis (data
15 preparation) using S language:

```
      all.data.princomp <- menuPrincomp(data =  
      by.compound.matrix, scores = T, cor = "Correlation",  
      na.action = T, print.short = T, print.importance = T,  
      print.loadings = T, cutoff.loadings = 0.1,  
20   plot.screplot = T, plot.loadings = T, plot.biplot = T,  
      plot.biplot.choices = c(1,2), predict.p = F)
```

 Example of clustering using a divisive hierarchical clustering
algorithm:

```
25       > div.hier.2.manhattan.cluster$call  
      diana(x = tmp.sum.by.comp, diss = F, metric =  
      "manhattan",  
          stand = T, save.x = T, save.diss = T)
```

30 Another embodiment utilizes existing tools for biological sequence similarity searches, classification, and phylogenetic analysis. In a particular embodiment, numbers in a numerical descriptor can be substituted by one or more of nucleic acid or amino acid codes according to a one of several sets of rules. Once

converted into a corresponding nucleotide or amino acid sequence representation, the fingerprints can be analyzed and compared using software and algorithms known in the art for genetic and peptide sequence comparisons, such as GCG, a product of Genetics Computer Group, with company headquarters in Madison WI. Select
5 embodiments comprising such approaches enable the use of a broad array of sophisticated algorithms to compare, analyze, and cluster gene and protein sequences. Many programs performing this task are known to those of ordinary skill in the art, such as for example, the PHYLIP (PHYlogeny Interference Package) a package of programs for inferring phylogenies (evolutionary trees) described in (Feldenstein, J.
10 1996 Methods Enzymol 266:418-427 and Feldenstein, J. 1981 J. Mol. Evol. 17(6):368-376).

Embodiments can perform such analysis based upon factors such as numerical value, statistical properties, relationships with other values, and the like. Further details of a step of manipulation are noted more particular below.

15 Fig. 7B illustrates a representative block flow diagram of simplified process steps for determining one or more descriptors of a state in the portions of the cells in the presence of the manipulation of step 704 of Fig. 7A in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill
20 in the art would recognize other variations, modifications, and alternatives. In a step 712, an image of a cell portion is obtained. In some embodiments, the cell portion is visualized with a fluorescently labeled marker that is specific for the portion or portions of interest. A cell portion can include, for example, one or more of the following: nuclei, Golgi apparatus, and other features. The cell portion may vary in
25 select embodiments according to the invention. Then, in a step 714, a digitized representation of the image obtained in step 712 is determined. In some embodiments, steps 714 and step 712 can comprise a single step. These embodiments use a digital imaging means such as a digital camera, to obtain a digital image of the target directly. Next, in a step 716, the digital representation of the image is
30 processed to obtain image features. Image features can include such quantities as area, perimeter, dimensions, intensity, aspect ratios, and the like. Then, in a step 718 descriptors can be determined from the image features. Descriptors can comprise scalar or vector quantities and can comprise the image features themselves, as well as

composed features, such as shape factor derived by a relationship $4\pi * \text{area} / \text{perimeter}$, and the like. Descriptors can also comprise statistical quantities relating to feature characteristics across a population of cells, such as a standard deviation, and average, and the like.

5 In a preferred embodiment, cells can be placed onto a microscope, such as a Zeiss microscope, or its equivalent as known in the art. A starting point, named Site A01, is identified to the microscope. A plurality of exposure parameters can be optimized for automated image collection and analysis. The microscope can automatically move to a new well, automatically focus, collect one or more images, at
10 one or more wavelengths, move to a next well, and repeat this process for all designated wells in a multiple well plate and for multiple plates. A file having a size and an intensity distribution measurement for each color and rank for each well can then be created for the images acquired. Based on this information, a user or a computer can revisit sites of interest to collect more data, if desired, or to verify
15 automated analysis. In a presently preferred embodiment, image automatic focus and acquisition can be done using computer software controlling the internal Z-motor of the microscope. Images are taken using a 10x, 20x, or 40x air long working distance objectives. Sometimes multiple images are collected per well. Image exposure times can be optimized for each fluorescent marker and cell line. The same exposure time
20 can be used for each cell line and fluorescent marker to acquire data.

Fig. 7C illustrates a representative block flow diagram of simplified process steps for obtaining images of cell portions of step 712 of Fig. 7B in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill
25 in the art would recognize other variations, modifications, and alternatives. The method is generally outlined by the steps below:

(1). In a step 720, a sample is provided to the imaging device. Samples can be provided in 96 well plates and the like. The sample may be loaded into a microscope, such as a Zeiss microscope or equivalent.

30 (2). In a step 722, a set of optical filters is selected to shine light of the appropriate wavelength to illuminate the first sample, which may be contained in a first well designated A01.

(3). In a step 724, an automatic focusing procedure is performed for the site. In a particular embodiment, the internal z-motor of the microscope which is attached to the objective nosepiece is used for automatic focusing of the microscope. In an alternative embodiments, the plate holding the samples is moved to perform
5 automatic focusing of the microscope, or focusing can be performed by moving optical components attached to the microscope and the like.

(4). In a step 726, images are collected for the site. Images can be collected for every color at every site. Present embodiments can provide images for up to four colors. However, embodiments are contemplated that can provide more
10 colors by using either a monochromator coupled with excitation filters which are on a filter wheel, or by digitally separating overlapping fluorophores. Those knowledgeable in the field will know that given calibration images of single fluorophores, a look-up table can be devised which will allow for the digital removal of fluorescence bleed-through of fluorescence which may occur in optical channels
15 other than the one for which that filter has been optimized in instances of using more than one fluorophore at once. Cell growth and density information is also collected. Cell density is determined by what percentage of the area being imaged is inhabited by cells. In some embodiments, imaging can be facilitated using one or more biosensors, molecules such as non-proteins, i.e., lipids and the like, that are
20 luminescently tagged. However, some embodiments can also use fluorescence polarization and the like. Fluorescence polarization is a homogeneous fluorescence technology where the excited state of the molecule lasts much longer than in normal fluorescence, taking seconds to minutes to reach equilibrium, obliterating the need to wash away fluorescence markers that are not specifically bound to a marker. Further,
25 embodiments can detect differences in spectral shifts of luminescent markers. Some fluorescence markers, such as Nile Red sold by Molecular Probes of Eugene, OR, will change its emission peak wavelength depending on its environment. One can detect these changes by monitoring the level of fluorescence at both wavelengths and reading out at ratio of the two.

30 (5). In a step 728, a determination is made whether more fields of view need to be taken for a particular color. If this is so, then processing continues at step 726 at a new site. Otherwise, processing continues with a decisional step 730.

Images can now be taken by repeating step 726. In a preferred embodiment 4 to 9 images are collected at each site.

(5). In a step 730, a determination is made whether more optical configurations need to be taken in order to obtain images for all differently-marked cell portions the sample. If this is so, then in a step 732 a new optical configuration is determined. Images for the new optical configuration can now be taken by repeating steps 726 and 728.

(6). In a decisional step 734, after all optical configurations and images for fields of view in a sample have been obtained, a determination is made whether any further samples remain to be analyzed. If so, a new sample is brought into view and processing continues with step 720. Otherwise, image processing is complete. In a presently preferable embodiment, image data can be stored on a CD ROM using a CD ROM burner, such as CRW4416 made by Yamaha of Japan. However, other mass storage media can also be used.

Fig. 7D illustrates a representative block flow diagram of simplified process steps for processing digitized representations of step 716 of Fig. 7B in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. The method is generally outlined by the steps below:

(1). In a step 740, a digitized image input is preprocessed .

Preprocessing might include, but is not limited to, such operations as background subtraction, thresholding, smoothing, adoptive filtering, edge enhancements, contrast enhancements, histogram equalization. A particular combination of preprocessing steps can be applied to images in successive steps or in parallel to copies of the image.

A simplified example of a smoothing and background subtraction procedure in a MatLab language is presented in computer code below:

```
function Isubtracted = cmBackgrSubtrl(I,k)
```

```
% cmBackgrSubtrl(I,k) - simple flat background (=modal*k)  
subtraction
```

```
% Y = cmBackgrSubtrl(I, k) - image Y is generated by
```

```

    % subtraction (with saturation) of modal pixel value of I
    multiplied by k
    % DEFAULT - k=1
    %
5   if (nargin == 1)
        k=1;
    end
    if (size(k)~=1)
        error('cmBackgrSubtrl: parameter k should be a number.
10  Exiting...');
    end

    %modpixnum = floor(size(I(:),1)/2);
    %sortedval = sort( double(I(:)) );
15  %modpixel = sortedval(modpixnum);
    modpixel = median(double(I(:)));
    bg = k*modpixel;

    Isubtracted = mmsubm( uint8(I), uint8(round(ones(
20  size(I))*k*modpixel )) );

```

An example of a procedure for thresholding in computer code (MatLab) is presented below:

```

function thresh = GetThreshByPerim1(I, M)
25  % GetThreshByPerim1(I) Finds optimal thresholding value
    for image I
    % N = GetThreshByPerim1(I) Finds thresholding value N for
    image I
    % N = GetThreshByPerim1(I, M) - tests threshold values up
30  to M
    % DEFAULT M = maximum pixel value in I
    % note that GetThreshByArea is significantly faster
    % finds a threshold value that causes the maximal change
    in the

```

```
% total perimeter of the objects (Russ ????)
% see Matlab_Auto_threshold1_1-23-99.doc for more details
% Note: works somewhat better on SMOOTH images (i.e.
medfilt2(I, [3 3]) two times

5
if (nargin == 0)
    error (strcat( mfilename, ' : at least one parameter
required'));
elseif (nargin == 1)
10    M = double(max(I(:)));      %test thresholds up to
maximum pixel value in I
elseif (nargin > 2)
    error (strcat (mfilename, ' : too many parameters'));
end

15
if (size(M)>1)
    error (strcat(mfilename, ' : argument M should be a
number'));
end

20
Minval = double( min(I(:)));
step = 1;

%generate vertical vector perims with total perimeters of
25 objects at different
%threshold values
for i=Minval : step : M
    bwI = im2bw(I, i/255);
    prI = bwperim(bwI);
30    pr = sum(prI(:));
    if (exist('perims', 'var') == 0) %perims is yet
undefined
        perims = pr;
    else
```



```

        perims = cat(1, perims, pr);
    end
end

5  % vector prdiffs contains differences between successive
   perimeters
   prdiffs = diff(perims);
   mindecrease = min(prdiffs);
   minvalues = find(prdiffs == mindecrease);
10  index_of_mindecrease = minvalues(1);
   thresh = index_of_mindecrease + 1;

   % =====end GetThresh1=====

```

15 Thresholding provides a specific intensity, such that pixels darker than the threshold are deemed black, and pixels lighter than the threshold are considered white. The thresholded image can be processed using binary image processing techniques in order to extract regions.

(2). In a step 742+744, the digitized image input is subjected to object
 20 identification. This can be accomplished by a variety of procedures, for example by thresholding or edge detection and subsequent morphological opening and closing. Edge detection can be accomplished by means of gradient-based or zero-crossing methods, such as Sobel, Canny, Laplassian, Perwitt, and other methods.

An example of object identification procedure based on Canny edge
 25 detection (in MatLab language) is presented below:

```

function Imask = cmMaskDNA1(I);
% cmMaskDNA1 - generates binary mask for cell nuclei
% through edge detection
30 % Imask = cmMaskDNA1(I)
% PARAMETERS
%   I - intensity image (grayscale)
% OUTPUT
%   Imask - BW image with objects from I

```

```

%
% For more details see Notebook Matlab_DNA_masking1_1-22-
99.doc
% Uses SDC Morphology Toolbox V0.7
5
if (nargin ~= 1)
    error('Wrong number of input parameters');
end
if (nargout ~= 1)
10    error('Wrong number of output parameters: one output
argument should be provided');
end

15  Imask = edge(I, 'canny');
    Imask = mm dil(Imask, mmsecross(1));
    Imask = mmero ( mmc lohole(Imask, mmsecross(1)));
    Imask = mmedgeoff(Imask, mmsecross(1));
    % note that mmedgeoff this command removed FILLED OBJECTS
20  but not touching OUTLINES.
    % these outlines can be removed by filtering:
    Imask = medfilt2(Imask, [5 5]);

    %=====end cmMaskDNA1
25  =====

```

However, embodiments can also use other techniques, such as Fast Fourier Transforms (FFT) and the like as known in the art without departing from the scope of the present invention.

30 (3). In a step 746, a plurality of region features can be determined. For example, in a representative embodiment, image features can include such quantities as area, perimeter, dimensions, intensity, aspect ratios, and the like. Features not directly related to individual objects are also being extracted.

An example of a procedure for extraction of some of the features (MatLab language) is presented below:

```

function OData = cmGetObjectsData(I, Ilabel)
5  % cmGetObjectsData returns array measurements of objects
  in image "I" masked by "Ilabel"
  % EV 2-3-99; 2-10-99
  % OData = cmGetObjectsData(I, Ilabel) returns an array of
  morphological and intensity measurements
10 %   taken from a grayscale image "I". Objects are
  identified on a mask image Ilabel, usually
  %   created by bwlabel()
  % OUTPUT:
  % Each row in the output array OData represents
15 individual object
  % columns contain the following measurements:
  %
  %   1 - Index ("number" of an object);      8 -
  Solidity;
20 %   2 - X coordinate of the center of mass; 9 - Extent;
  %   3 - Y coordinate      "-"      ; 10 - Total
  Intensity;
  %   4 - Total Area (in pixels);              11 - Avg.
  Intensity;
25 %   5 - Ratio of MajorAxis/MinorAxis;        12 - Median
  Intensity;
  %   6 - Eccentricity;                        13 - Intensity of
  20% bright pixel
  %   7 - EquivDiameter;                       14 - Intensity of
30 80% bright pixel
  %
  % For details on morphological parameters see information
  on MatLab imfeature();

```

```
% Intensity parameters are either obvious or are
documented in comments in this file.

if (nargin ~= 2)
5   error ('function requires exactly 2 parameters');
end
if (nargout ~= 1)
    error ('function has 1 output argument (array X by
14) ');
10 end

% finished checking arguments

% first collect morphological parameters in a structure
15 array:
ImStats = imfeature(Ilabel, 'Area', 'Centroid',
    'MajorAxisLength',...
    'MinorAxisLength', 'Eccentricity', 'EquivDiameter',
    ...
20 'Solidity', 'Extent', 8 );

% now convert it into array (matrix) while collecting
intensity data for each object:

25 %preallocate output array:
numobjects = size(ImStats, 1);
OData = zeros(numobjects, 14);
%now convert ImStats into array and add intensity data to
it
30 for k=1:numobjects
    OData(k, 1) = k;
    OData(k, 2) = ImStats(k).Centroid(1);
    OData(k, 3) = ImStats(k).Centroid(2);
    OData(k, 4) = ImStats(k).Area;
```

```

        OData(k, 5) = (ImStats(k).MajorAxisLength) /
        (ImStats(k).MinorAxisLength);
        OData(k, 6) = ImStats(k).Eccentricity ;
        OData(k, 7) = ImStats(k).EquivDiameter;
5       OData(k, 8) = ImStats(k).Solidity;
        OData(k, 9) = ImStats(k).Extent;

        % now collect and assign intensity parameters from
        image I
10
        object_pixels = find( Ilabel == k);
        object_area = size(object_pixels, 1); %same as total
        number of pixels in the object
        object_intensities = double(I(object_pixels)); %
15    need to convert to double to do math
        sorted_intensities = sort(object_intensities); %
        will need to get median, 20% and 80% pixels
        total_intensity = sum(object_intensities, 1);
        avg_intensity = total_intensity / object_area;
20    median_intensity = sorted_intensities( floor(
        object_area/2 ) + 1 );
        pix20 = sorted_intensities( floor(object_area*0.2)+1
        ) ; %brightest pixel among dimmest 20%
        pix80 = sorted_intensities( floor(object_area*0.8)+1
25    ) ;

        OData(k, 10) = total_intensity;
        OData(k, 11) = avg_intensity;
        OData(k, 12) = median_intensity;
30    OData(k, 13) = pix20; %brightest pixel among dimmest
        20%
        OData(k, 14) = pix80; %dimmest pixel among brightest
        20%
        end %for

```

```
%===== end function
cmGetObjectsData() =====
```

- 5 (4). In a step 748, quantitative descriptors, characterizing cell state are calculated based on the feature measurements extracted at step 746. For example, histogram distribution of intensities of cell nuclei provides information about the population cell cycle stages.

In a particular embodiment according to the present invention, data analysis techniques for describing the fluorescence patterns of cell portions in multiple cell lines in the presence and absence of compounds are provided. Automated image analysis techniques can include determining one or more regions from around nuclei, individual cells, organelles, and the like, called "objects" using a thresholding function. Objects that reside on the edge of an image can be included or excluded in various embodiments. An average population information about an object can be determined and recorded into a database, which can comprise a database text file or Excel spreadsheet, for example. However, embodiments can use any recording means without departing from the scope of the present invention. Values measured can be compared to the visual image. One or more types of numerical descriptors can be generated from the values. For example, descriptors such as a number of objects, an average, a standard deviation of objects, a histogram (number or percentage of objects per bin, average, standard deviation), and the like can be determined.

In a particular embodiment according to the present invention, data can be analyzed using morphometric values derived from any of a plurality of techniques commonly known in the art. For example, a software package called MetaMorph Imaging System, provided by Universal Imaging Corporation, a company with headquarters in West Chester, PA and NIH Image, provided by Scion Corporation, a company with headquarters in Frederick, Maryland.

30 Fluorescent images can be described by numerical values, such as for example, an area, a fluorescence intensity, a population count, a radial dispersion, a perimeter, a length, and the like. Further, other values can be derived from such measurements. For example, a shape factor can be derived according to a relationship

4π * area / perimeter. Other values can be used in various embodiments according to the present invention. Such values can be analyzed as average values and frequency distributions from a population of individual cells.

In a particular embodiment according to the present invention, techniques for the automatic identification of mitotic cells are provided. Image analysis techniques employing techniques such as multidimensional representations, frequency-based representations, multidimensional cluster analysis techniques and the like can be included in various embodiments without departing from the scope of the present invention. Techniques for performing such analyses are known in the art and include those embodied in MatLab software, produced by MathWorks, a company with headquarters in Natick, MA.

Scalar values providing efficacious descriptors of cell images can be identified using the techniques of the present invention to perform predictive analysis of drug behavior. In a presently preferred embodiment, a plurality of heterogeneous scalar values can be combined to provide descriptors for each manipulation. By applying predictive analysis routines to the collections of these descriptors, predictive information about any number of manipulations and cell interactions can be extracted.

Fig. 7E illustrates a representative block flow diagram of simplified process steps for analyzing image feature values to obtain descriptors of cell state of step 718 of Fig. 7B in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. Fig. 7E illustrates an input data of descriptors of known manipulations 319. A step 320 of reformatting and transforming data 319 to formats suitable for analysis is performed. Additionally, a "cleaning" process can eliminate outlying data points and the like in the data. Then, in a step 322, a decision is made whether to continue with step 324 or with step 326 based upon determining a particular type of analysis appropriate for the present application or particular type of prediction. If decisional step 322 determines processing should continue with step 324, then, in that step, an error estimate using a set of test descriptors is performed to estimate the quality of a prediction and processing continues with step 320. Once an optimal prediction is achieved, processing continues with step 326. In step 326, optimal transformation parameters and prediction methods are selected for use in

steps 328 and 330 which analyze data about an unknown manipulation. In a step 328, a solution is generated based upon any of techniques including training a neural network, solving a mathematical equation, applying decision tree rules and/or the like. In a step 330, an input data set of unknown descriptors 318 is reformatted and
5 transformed based upon the optimal transformation parameters selected in step 326 using the transformation procedures in steps 320, 322 and 324. In a step 332, predictions techniques are applied to the reformatted manipulations from step 330 and the solution generated in step 328 and a plurality of properties of known manipulations 317 (e.g., therapeutic properties, and the like) in order to determine a
10 prediction of properties of unknown manipulation 316.

Fig. 7F illustrates a representative block flow diagram of simplified process steps for a method of mapping a manipulation of cells to a physiological characteristic in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein.
15 One of ordinary skill in the art would recognize other variations, modifications, and alternatives. The method is generally outlined by the steps below:

(1) In a step 750, a plurality of cells, e.g., dead, live, cell fractions or mixtures of cells are provided.

(2) Then, in a step 752, the plurality of cells is manipulated, where
20 manipulation occurs using a source(s) from one or a combination selected from an electromagnetic, electrical, chemical, thermal, gravitational, nuclear, temporal, or a biological source.

(3) Next, in a step 754, a feature value is captured from the plurality of cells. The feature value can include one or any combination of characteristics such as
25 cell count, area, perimeter, length, breadth, fiber length, fiber breadth, shape factor, elliptical form factor, inner radius, outer radius, mean radius, equivalent radius, equivalent sphere volume, equivalent prolate volume, equivalent oblate volume, equivalent sphere surface area, average intensity, total intensity, and optical density. This list is not meant to be limiting.

(4) Then, in a step 756, a degree of presence of one or more feature values is assigned for each manipulation.
30

(5) In a step 758, the feature values from the plurality of cells are stored in memory locations. From the memory locations the values can be used for

statistical analyses to produce predictive information about the relatedness of the descriptors of the manipulations to one another. This information is used to infer properties of the manipulations.

Fig. 7G illustrates a representative block flow diagram of a simplified process steps for a method for populating a database with manipulated biological cell information in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. The method is generally outlined by the steps below:

10 (1) In a step 760, a plurality of cells in various stages of the cell cycle, A montage image that was used as a source to generate data in Appendix A is presented in Fig. 12., such as for example, the stages of interphase, prophase, metaphase, anaphase, and telophase are provided.

(2) Then, in a step 762, each of the cells in the various stages of mitotic development is manipulated.

(3) Next, in a step 764, an image of the plurality of manipulated cells is captured using image acquisition techniques in order to provide a morphometric characteristic of each of the manipulated cells.

(4) As a preferable option, in a step 766, an image database may be populated with the image of the plurality of manipulated cells.

(5) Following step 764 or optional step 766, a morphological value is calculated from the image in a step 768.

(6) In a step 770, the database is populated with the morphological value.

25 Fig. 7H illustrates a representative block flow diagram of simplified process steps for a method for populating a database with manipulated biological information, e.g., image acquisition parameters, image feature summary information, and well experimental parameters in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. Fig. 7H illustrates a step 780 in which cells are placed into site on a plate and a manipulation is applied. Then, in a step 781 an image is taken of the cells. In step 782, the image is transferred to an image archive

30

database. Then, in a step 783, well experimental parameters are entered into the database 787. Well experimental parameters can include cell type, manipulation and the like. In a step 784, image acquisition parameters are transferred to database 787. Image acquisition parameters can include file name, fluorophores and the like. In a
5 step 785, the image acquired in step 781 is analyzed. Then, in step 786, an image feature summary from the analysis step 785 is transferred to database 787.

In step 788, a lookup table for all analyses is provided to database 787. The lookup table provides information about the analyses. In a step 789, a query of database 787 for process data is performed. The results are reformatted. Then in a
10 step 790, the database 787 is queried. Next, in a step 791, features of the manipulations stored in the database are combined and reduced. Next, in a step 793, reduced features of step 791 can be compared. In a step 792, the results of step 793 are recorded in database 787. Then, in a step 794, a report of predictions based on comparisons performed in step 793 is generated.

15 Fig. 7I illustrates a representative block flow diagram of simplified process steps for acquiring images of manipulated biological information, e.g., cells, cell tissues, and cell substituents in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations,
20 modifications, and alternatives. Fig. 7I illustrates a step 770 in which a user sets up an image analysis procedure. Then, in a step 772, an image is read into image analysis software. Next, in a step 774, patterns and objects are identified in the image using one or more algorithms. Next, in a step 776, sets of features are extracted from the image. Then, in a step 778, feature information, descriptor values and the like are
25 exported to the database, such as database 787 of Fig. 7H, for recording. Next, in a decisional step 779, a determination is made whether any more images should be taken. If this is so, processing continues with step 772. Otherwise, image acquisition processing is completed.

Fig. 7J illustrates a representative block flow diagram of simplified
30 process steps for populating, acquiring and analyzing images of manipulated biological information in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations,

modifications, and alternatives. Fig. 7J illustrates a step 300 of placing a plate onto an imaging stage and reading a bar code. Then, in a step 301 an autofocus procedure is performed. Next, in a step 302, a first optical filter configuration is selected and an image is collected. Then, in a decisional step 303, a determination is made whether
5 more than one image per optical configuration can be taken. If so, then, in a step 304, a new position within the well is targeted and another image is collected. Then, in a decisional step 305, a determination is made whether any more images need to be collected. If this is so, step 304 is repeated until all images for a particular well have been collected. After one or more images are collected for the well, in a step 306, the
10 stage is returned to a starting position within the well, and a montage is created from collected images. The results are named with a unique file name and stored.

In a decisional step 307, a determination is made whether any more optical channels in the well can be imaged. If this is so, then in a step 308 the next optical filter configuration is selected and an image is collected. Processing then
15 continues with decisional step 303, as described above. Otherwise, if no further optical channels in the well can be imaged, then in a decisional step 309 a determination is made whether any wells remain to be imaged. If not all wells have been imaged, then in a step 310, the stage moves to the next well and processing continues with step 301, as described above. Otherwise, if all wells on the plate have
20 been imaged, then in a decisional step 311, a determination is made whether any more plates can be processed. If this is so, then processing continues with step 300 as described above. Otherwise, in a step 312, the information is stored to a CD or other storage device as a backup.

Fig. 7K illustrates a representative block flow diagram of simplified
25 process steps compound based upon information about effects of one or more known compounds on a cell population in a particular embodiment according to the present invention. This diagram is merely an illustration and should not limit the scope of the claims herein. One of ordinary skill in the art would recognize other variations, modifications, and alternatives. Fig. 7K illustrates a step 340 of populating a database
30 with descriptors for known compounds. Such descriptors can be determined from imaging the cell population. However, in some embodiments, descriptors can be derived by measurements and combinations of measurements and the like. Then, in a step 342, descriptors for the unknown compound are determined from imaging a

second cell population. The second cell population has been treated with the unknown compound. Then, in a step 344, a relationship between the descriptors determined from the unknown compound with the descriptors determined from the known compounds can be determined. Finally, in a step 346, an inference can be made about the unknown compound based upon the descriptors of the known compounds from the relationship determined in step 344.

Accordingly, the present invention provides a novel database design. In a particular embodiment according to the present invention, a method for providing a database comprises measurement of a potentially large number of features of one or more sub-cellular morphometric markers. Markers can be from any of a large variety of normal and transformed cell lines from sources such as for example, human beings, fungi, or other species. The markers can be chosen to cover many areas of cell biology, such as, for example markers comprising the cytoskeleton of a cell. The cytoskeleton is one of a plurality of components that determine a cell's architecture, or "cytoarchitecture". A cytoarchitecture comprises structures that can mediate most cellular processes, such as cell growth and division, for example. Because the cytoskeleton is a dynamic structure, it provides a constant indication of the processes occurring within the cell. The cytoarchitecture of a cell can be quantified to produce a one or more scalar values corresponding to many possible cellular markers, such as cytoskeleton, organelles, signaling molecules, adhesion molecules and the like. Such quantification can be performed in the presence and absence of drugs, peptides, proteins, anti-sense oligonucleotides, antibodies, genetic alterations and the like. Scalar values obtained from such quantification can provide information about the shape and metabolic state of the cell.

In a presently preferred embodiment, scalar values can comprise morphometric, frequency, multi-dimensional parameters and the like, extracted from one or more fluorescence images taken from a number of cellular markers from a population of cells. Two or more such scalar values extracted from a plurality of cell lines and markers grown in the same condition together comprise a unique "fingerprint" or descriptor that can be incorporated into a database. Such cellular descriptors will change in the presence of drugs, peptides, proteins, antisense oligonucleotides, antibodies or genetic alterations. Such changes can be sufficiently unique to permit a correlation to be drawn between similar descriptors. Such

correlations can predict similar properties or characteristics with regard to mechanism of action, toxicity, animal model effectiveness, clinical trial effectiveness, patient responses and the like. In a presently preferred embodiment, a database can be built from a plurality of such descriptors from different cell lines, cellular markers, and
5 compounds having known mechanisms of action (or structure, or gene response, or toxicity).

The present invention also provides database and descriptor comparisons according to other embodiments. In a particular embodiment according to the present invention, measurement of scalar values or features can provide
10 predictive information. A database can be provided having one or more "cellular fingerprints" comprised of descriptors of cell substance interactions of drugs having known mechanisms of action with cells. Such descriptors can be compared using a plurality of techniques, such as a technique of creating "phylogenetic trees" of a statistical similarity between the descriptors from various drugs. In a present
15 embodiment, scalar, numeric values can be converted into a nucleotide or amino acid letter. Once converted into a corresponding nucleotide representation, the descriptors can be analyzed and compared using software and algorithms known in the art for genetic and peptide sequence comparisons, such as GCG, a product of Genetics Computer Group, with company headquarters in Madison WI. In an alternative
20 embodiment, numeric values for the fingerprints can be used by comparison techniques. A phylogenetic tree can be created that illustrates a statistical significance of the similarity between descriptors for the drugs in the database. Because the drugs used to build the initial database are of known mechanism, it can be determined whether a particular scalar value in a descriptor is statistically predictive. Finally, a
25 compound fingerprint with no known mechanism of action can be queried against the database and be statistically compared and classified among the drugs in the database that the compound most resembles.

In a particular embodiment, relationships between measured morphometric properties and features of images and physiological conditions can be
30 determined. Relationships can include, for example, treatment of different cell lines with chemical compounds, or comparing cells from a patient with control cells, and the like. In a presently preferable embodiment, a clustering can be performed on acquired image descriptors. Some embodiments can comprise statistical and neural

network - based approaches to perform clustering and comparisons of various descriptors. The foregoing is provided as merely an example, and is not intended to limit the scope of the present invention. Other techniques can be included for different types of data. In some embodiments, clustering and comparing can be performed on features extracted from cell images. In a presently preferable embodiment, procedures for comparisons and phylogenetic analysis of biological sequences can be applied to data obtained from imaging cells.

Select embodiments comprising such approaches enable the use of a broad array of sophisticated algorithms to compare, analyze, and cluster gene and protein sequences. Many programs performing this task are known to those of ordinary skill in the art, such as for example, the program Phylip, available at <http://evolution.genetics.washington.edu/phylip.html>, and other packages listed at <http://evolution.genetics.washington.edu/phylip/software.html>. However, select embodiments according to the present invention can comprise a technique of statistical classification, statistical clustering, distance based clustering, linear and non-linear regression analysis, self-organizing networks, and rule-based classification.

Embodiments can perform such analysis based upon factors such as numerical value, statistical properties, relationships with other values, and the like. In a particular embodiment, numbers in a numerical descriptor can be substituted by one or more of nucleic acid or amino acid codes. Resulting "pseudo-sequences" can be subjected to analysis by a sequence comparison and clustering program.

Other types of databases can also be provided according to other embodiments. The database includes details about the properties of a plurality of standard drugs. When the descriptor of a test compound is compared to the database, predictions about the properties of the test compound can be made using any known property of the other compounds in the database. For example, properties about a compound in the database could include structure, mechanism of action, clinical side effects, toxicity, specificity, gene expression, affinity, pharmacokinetics, and the like. The descriptor of a compound of unknown structure from a natural products library could be compared to the descriptors of compounds with known structure and the structure could be deduced from such a comparison. Similarly, such information could lead to better approaches to drug discovery research including target validation

and compound analogizing, as well as pre-clinical animal modeling, clinical trial design, side effects, dose escalation, patient population and the like.

According to the present invention, databases can be integrated with and complementary to existing genomic databases. Differential genomic expression strategies can be used for drug discovery using database technology. In one particular embodiment, cell data and cellular response data can be associated with a genetic expression profile assay to form a single assay. Live cells expressing fluorescence markers can be treated with a drug, imaged and analyzed for morphometry; and then analyzed for mRNA for expression. Such embodiments can provide rapid development of tools to link cellular behavior with functional genomics.

Database methods according to the present invention can be used to predict gene function and to assist in target validation. Databases that include genetic diversity, i.e., having cellular descriptors from cells of differing genetic backgrounds (tumor, tissue specific, and gene knock out cell lines), can provide the capability to compare cells of unknown genetic background to those in the database. Similarly, the descriptor of an unknown cellular portion in the presence of multiple drugs can be queried against the descriptors of the known markers in the database. For example, if an unknown gene is tagged with Green Fluorescent Protein (GFP), the database may be used to identify the cellular portions for which that unknown gene encodes.

According to the present invention, target validation and specialized cell-based assay screening can be performed using database systems and methods to serve as a universal high-throughput cell-based assay that can evaluate the molecular mechanism of drug action. As new genes are isolated and identified, a large collection of available gene-based knowledge is becoming available. From this large collection of new genes, potential protein targets can be identified using the genomic tools of sequence analysis and expression profiling. However, unless a gene mutation is tightly linked to a disease state, further validation of individual targets is a time consuming process, becoming a bottleneck in drug discovery. Furthermore, robotics and miniaturization are making "High Throughput Screening (HTS)" the industry standard, substantially reducing the time and cost of running a target-based biochemical assay. Therefore, it is now possible to routinely screen large libraries and use a resulting "hit" to validate the target. In such approaches, a specialized cell-based assay would be developed to test hits for each target. Since this often involves

the creation of cell lines expressing new markers, this stage may also become a bottleneck that cannot keep pace with HTS. In addition, these cell-based assays may not be amenable to high-throughput screening, making it difficult to test the increasing number of analogs arising from combinatorial chemistry.

5 In a particular embodiment according to the invention, a rapid characterization of large compound libraries for potential use as pharmaceutical products can be provided by predicting properties of compounds that relate to the compounds' potential as bioactive drugs. In many drug discovery situations, virtually millions of compounds can be passed through a HTS assay against a small number of
10 validated targets. These assays produce hundreds to thousands of potential hits. These hits can then be subsequently screened by a pipeline of secondary and tertiary screens to further characterize their specificity, often time completely missing non-specific interactions with other proteins. Techniques according to the present invention can provide a replacement to such screening operations by providing
15 information about cellular accessibility and mechanism of action for the hits coming from a HTS system. Furthermore, it can replace the biochemical HTS assay and allow rapid and accurate identification of attractive compounds from large libraries without an intervening biochemical assay. The cell information can be predictive of whether to continue into an animal model for each compound, and which animal model to
20 pursue.

The principles of the present specifically contemplate a wide variety of research methodologies, or usage scenarios, implementing these principles. The following discussion of three such scenarios is by way of illustration and not limitation. Study of the principles enumerated herein will render evident to those
25 skilled in the art certain additional methodologies or usage scenarios enabled by the teachings hereof. The present invention specifically contemplates all such modifications. The following description presents some specific embodiments and scenarios that represent a broader use of cellular phenotypic data and characterizations to deduce mechanisms of action and other features of cellular
30 responses to various stimuli. Such procedures generally involve producing a quantitative cellular phenotype based upon two or more cellular attributes and then comparing that phenotype to phenotypes previously stored and indexed. Such

procedures make use of databases or other repositories of biological information. The invention is not limited to the specific embodiments described here.

Considering first the procedure 2000 depicted in Figure 20, a compound has been identified as having a particular cellular activity. See 2004. For example, a compound may be found to inhibit the growth of certain cancer cell *in vitro* by a specific and desired mechanism of action. This may be a particular company's "gold standard."

Next, the compound is analyzed at 2006 in terms of its effect on one or more cell lines. More specifically, the compound is linked, virtually, to a particular phenotype. Two or more values or measures of cellular attributes characterize that phenotype. These attributes are quantified in the context of specific cellular markers.

In one example, the cellular marker is an organelle such as a nucleus or Golgi apparatus. Measured attributes useful for characterizing an associated phenotype include geometric parameters (e.g., size, shape, and/or location of the organelle) and composition (e.g., concentration of particular biomolecules within the organelle).

The phenotype may be characterized by administering the compound of interest to various cell lines and in various concentrations. In each example within this matrix, the attributes of interest are measured. Ultimately, certain phenotypic features (combinations of attribute values) are associated with the compound of interest. These features provide a template for the phenotype.

Next, using the phenotype as identified at 2006, the process identifies other compounds providing similar features. The goal here is to present a list of compounds having a mechanism of action similar to that of the compound that started the process. This allows researchers to identify a mechanism of action, if not already known, for their compound and to draw conclusions based upon their compound's link to other known compounds (which may not be chemically/structurally similar to the compound of interest).

Identifying similar compounds based upon phenotype can take many paths. Most will involve some mathematical basis. For example, the phenotype defined at 2006 can be represented as a fingerprint or vector comprised of multiple scalar values of cellular attributes (as described above). The phenotype representation can then be compared against known phenotypes characterized by the same format

(e.g., they are all characterized as vectors having the same attribute set, but with different values of the attributes). The comparison may be as simple as a Euclidean distance or more sophisticated as a neural network or multivariate statistical correlation.

5 The known compounds and associated phenotypes may be stored as database records or other data structures that can be queried or otherwise accessed as part of the identification procedure. The compounds may also be associated with other relevant data such as clinical toxicity, cellular toxicity, hypersensitivity, mechanism of action, etc. (when available).

10 Compounds found to be sufficiently similar to the starting compound are returned for consideration by researchers. A data processing system may rank such compounds based on degree of similarity to the starting compound. In some cases, the system may even provide similarity scores associated with the listed compounds.

15 Often researchers wish to determine whether their particular compound has clinical or biochemical effects beyond those that they are already aware of. In a typical scenario, the compound of interest was selected based upon its strong binding a target or its stimulation or inhibition of cell growth in a particular cell line. The process associated with 2010 has likely identified the compound of interest as having
20 a particular mechanism of action based on phenotypic similarity to other compounds having a similar mechanism of action. However, within the region of biochemical space, there may be subspaces (characterized by subphenotypes) that correspond to separate properties. For example, within the phenotypic space associated with one mechanism of action, there may be subspaces associated with clinical toxicity,
25 cellular toxicity (likely overlapping the clinical toxicity space), and little or no toxicity. Obviously, a researcher would like to know whether her compound is likely to be toxic.

 Thus, the process 2000 may include characterizing the compound of interest in terms of its distance from (i.e., similarity to) specific phenotypes having
30 known characteristics. In a typical example, the known characteristic is toxicity. This feature allows the researcher to quantify her compound in terms of mechanism of action AND toxicity (or in terms of two or more other relevant properties associated

with phenotype). To allow simple ranking or characterization, compounds of interest may be scored according to a simple or weighted Boolean expression.

A second scenario of interest is depicted in Figure 21. This scenario again defines a phenotype in terms of a quantifiable vector or other measure.

5 However, rather than using a compound of interest to generate the phenotype, some other cellular stimulus is used to generate the phenotype.

As shown, a process 2100 begins with receipt of cells of interest. See 2104. In many situations, the cells are produced by a genetic or epigenetic process that affects the expression level or activity of a particular protein. More generally,
10 any cellular stimulus (e.g., radiation level and type, gravity level, magnetic field, acoustic perturbations, etc.) can be used to generate the cell line of interest. Importantly, this stimulus affects the phenotype and can be correlated therewith.

In the context of drug discovery, a gene encoding for a particular target can be genetically knocked out, underexpressed, overexpressed, expressed in a non-
15 native state, etc. This may be accomplished via standard procedures involving genomic modification, translation or transcription apparatus modification (e.g., use of antisense nucleic acids), blocking target activity (using antibodies to a receptor site for example), and the like. These processes will generally affect the phenotype in some quantifiable way. Importantly, they clearly and unambiguously define a cellular
20 phenotype associated with altering the activity of the target protein.

At 2106, the process involves measuring one or more cellular features from the cell line of interest to define/quantify the phenotype. This may be accomplished as described above with reference to 2006. Next, at 2108, the cellular phenotype generated in this manner is used to identify and rank a set of compounds
25 associated with the phenotype. This operation may proceed in the manner of operations 2008 and/or 2010 from Figure 20.

Finally, at 2110, the process clusters the compounds returned at 2108 by a mechanism of action. The operation 2106 has tightly bound a mechanism of action to a phenotype. Various compounds characterized and stored in a system
30 database may be tentatively assigned a mechanism of action or may have no suggested mechanism of action. By matching their virtual phenotype to the phenotype generated at 2106, one can create or strengthen an association between the compounds and mechanism of action relevant to the stimulus at 2104.

Considering now Figure 22, a third scenario is depicted. This scenario again involves using a virtual phenotype to glean information relevant to a mechanism of action or other cellular activity. In this case, assay data from a group of compounds (e.g., a primary or focused library) is used to elucidate a phenotype.

5 As shown, a process 2200 begins by identifying a target protein. See 2204. Then, at 2206, the process involves identifying positive and negative biochemical hits. More generally, this may involve ranking a number of compounds based upon their interaction with the target. In a specific case, the compounds are ranked based upon their binding affinities to or ability to inhibit the enzymatic activity
10 of the target protein.

 After the compounds have been characterized in some manner based upon their interaction with the target, they are used to define a cellular phenotype. See 2208. Generally, the techniques to accomplish are the same as described with reference to operation 2006 of Figure 20. In this case however, one may obtain a
15 strong correlation between mechanism of action (involving the target) and phenotype by using multiple of the compounds identified at 2206. For example, some of the "best hits" may be administered to cell lines in various concentrations. And some of the least effective compounds may also be administered. Cellular attributes that are more strongly exhibited with increasing concentration of the best hits (and not
20 exhibited or exhibited only weakly upon administration of the negative hits) can be used to define the virtual phenotype. In a related approach, compounds having widely varying levels interaction with the target are administered to cells. Those cellular attributes that vary linearly or at least monotonically with the degree of interaction between the target and compound represent attributes that can be used to define the
25 virtual phenotype.

 After the cellular phenotype has been defined, previously characterized compounds may be clustered with that phenotype. See 2210. As with operation 2110 of Figure 2, this may create or strengthen an association between a mechanism of action and various compounds in a database.

30 Finally, and optionally, procedure 2200 may provide a "higher resolution" mechanism of action for the compounds identified at 2206. See 2212. Presumably interaction with the target suggests a specific mechanism of action or at least some aspect of a mechanism of action. However, a given target may participate

in a larger cellular mechanism of action – unknown to researchers. Further, a compound may that binds with the target may participate in multiple mechanisms of action – some of which do not involve the target. By linking the target (and its positive hits) to a particular phenotype, some of these additional cellular level activities can be elucidated. The defined phenotype may have been previously identified as associated with other mechanisms of action or higher resolution mechanisms of action. Thus, the phenotype identified at 2208 can be leveraged to generate a higher resolution mechanism of action at 2212.

As suggested in the above discussion, compounds and associated phenotypes may be stored as database records. Such databases can take on many flavors. In one example, a database includes various pieces of information relevant to oncology. Such database may include numerous compounds classified by cellular phenotype, mechanism of action, toxicity, etc. More specifically, the database may include data on commercially available compounds clustered by cellular phenotypes corresponding to mechanisms of action. Further the databases of interest may extended or combined (via standard relational tables and algebra for example) to include additional data such as pharmacology data, cellular genomics data, gene expression data, protein expression data, etc. In a specific example, the database includes measurements made on a subset of the NCI60 cell lines, using DNA, Golgi apparatus, and/or microtubules as markers for defining the phenotypes. Other data includes dosage response information, variation in effect over time, etc. The compounds populating the database could include known National Cancer Institute oncology study compounds. In a specific embodiment, the compound set includes some or all of the compounds mentioned in the article “A gene expression database for the molecular pharmacology of cancer,” Nature Genetics, 24, pp. 236-244 (March 2000).

Various biological analyses may be conducted to develop additional information for characterizing compound mechanisms of action, etc. For example, a cell count analysis may be used to develop dose response curves, GI 50 data, etc. The cell cycle may also be analyzed to find out how various stages in the cycle vary in response to particular stimuli. The Golgi apparatus may be analyzed to determine whether it is in a normal state, a dispersed state, a diffused state, etc. As another example, tubulin may be analyzed to determine whether it is normal, de-polymerized,

over-polymerized, bundled, etc. Obviously, combinations of such analyses may be performed. For example, properties of the Golgi apparatus or tubulin may be analyzed over one or more cell cycles.

In some embodiments, techniques according to the present invention
5 can provide tools for the later stages of drug development such as clinical trial design and patient management. The properties of known drugs, such as clinical trial and patient response information, will be used in a similar fashion as the pre-clinical information to provide predictions about the properties of novel compounds. Because the human cell is the locus of drug action, a database containing drug-cell interactions
10 will be able to provide predictive value for this aspect of drug development.

Although the above has generally been described in terms of specific hardware, software, and methods, it is understood that many alternatives can exist. In particular, the present invention is not limited to a particular kind of data about a cell, but can be applied to virtually any cellular data where an understanding about the
15 workings of the cell is desired. Thus, in some embodiments, the techniques of the present invention could provide information about many different types or groups of cells, substances, and genetic processes of all kinds. Of course, one of ordinary skill in the art would recognize other variations, modifications, and alternatives. Some examples according to the present invention are provided below.

20

EXPERIMENTS

To prove the principle and demonstrate the objects of the present invention, experiments have been performed to determine the effects of manipulations on cell structure using imaging and analysis techniques applied to a variety of
25 situations. These experiments were performed by growing multiple cell lines in the presence of multiple compounds, or substances. Cells were fixed and stained with fluorescent antibodies or labels to multiple cellular portions. One or more images of the cells were then obtained using a digital camera. Descriptors were built by quantifying and/or qualifying patterns of one or more feature from each image in the
30 cell lines under study. A database was built from the descriptors. As the database grows, it should be able to predict the mechanism of action of an unknown drug by comparing its effect with the effects of known compounds or to identify data clusters within large libraries of compounds.

In a first experiment, an automated method to count the number of cells and differentiate normal, mitotic, and apoptotic cells was created.

Approximately, 5,000 HeLa cells were plated per well in a 96 well plate and grown for 3.5 days. The cells were fixed with -20° MEOH for 5 minutes, washed with TBS for 15 minutes, and then incubated in 5 mg/ml Hoechst 33342 in TBS for 15 minutes. Then, 72 images were collected with a 40x objective and 75 ms exposure time.

The analysis was performed on objects that met a certain size criteria that was based on 1) measuring the size of objects in the image that were clearly not cells and 2) excluding the first peak of the area histogram (Fig. 8B values 1-4654).

Histograms of the individual object data were generated for each type of feature. Fig. 8A shows the histogram for average intensity, and Fig. 8B shows histogram data for the area of each object. Fig. 8C shows the scatter plot of the average intensity vs. the area of all of the objects. The pattern of the scatter plot showed an interesting pattern: a large cluster of cells in one region of the graph, with a scattering of object points in other regions. Because mitotic structures are identified as particularly bright objects, most likely due to the biological fact that the chromatin is condensed, the original Hoechst images could be used to identify which cells were either undergoing mitosis, or otherwise looked abnormal. Manual inspection of 917 cells resulted in the classification of each object. Fig. 8D shows a graph where each type of cellular classification is delimited. This graph clearly shows that the mitotic nuclei are brighter than the interphase nuclei. Further, the different phases of the cell cycle can be separated using these two features. Figs. 8E-8F show bar graphs of the average and standard deviations of the areas and average intensities for each cell classification type. These graphs show that interphase nuclei are statistically less bright than mitotic nuclei and that telophase nuclei are statistically smaller than other mitotic nuclei.

Each image was thresholded to an intensity level of 20. A standard area value was set at 9500 pixels. Automated information gathering about all of the objects was done and collected into an Excel spreadsheet (for more information see, section on imaging system). The following information was recorded:

IMAGE NAME
OBJECT #

AREA
STANDARD AREA COUNT
PERIMETER
FIBER LENGTH
FIBER BREADTH
SHAPE FACTOR
ELL. FORM FACTOR
INNER RADIUS
OUTER RADIUS
MEAN RADIUS
AVERAGE INTENSITY
TOTAL INTENSITY
OPTICAL DENSITY
RADIAL DISPERSION
TEXTURE DIFFERENCE MOMENT
EFA HARMONIC 2, SEMI-MAJOR AXIS
EFA HARMONIC 2, SEMI-MINOR AXIS
EFA HARMONIC 2, SEMI-MAJOR AXIS
ANGLE
EFA HARMONIC 2, ELLIPSE AREA
EFA HARMONIC 2, AXIAL RATIO
EFA HARMONIC 3, SEMI-MINOR AXIS

The following results were obtained:

- 1,250 objects were counted
- 201 of those objects has standard area counts > 2 (area > 19000 pixels)
- 195 objects had areas < 6000 pixels
- 1529 objects estimated in total
- 1328 object areas are > 6000 pixels
- The data was reduced to 917 objects that were $6000 < \text{area} < 19000$
- For the 917 objects a scatter plot of area vs. average intensity and a histogram of the average intensity were generated.

- 116 objects that had average intensity intensities > 60 were manually looked at to determine their morphology.

- Of those 116 objects:

6 were dead or indistinguishable

4 were interphase

30 were prophase

32 were metaphase

24 were anaphase

20 were telophase (10 pairs)

- 12 prophase objects were missed because of gray scale cut off. (8 of those prophase cells had gray scale values > 57 , as did 7 interphase)
- 1 telophase object was missed because it was too small (< 6000)
- 1 prophase object was missed because it was too big (> 1900)
- 16 mitotic objects were missed because they were parts of objects with standard count > 2 .

In sum, out of 917 single objects, the analysis correctly identified 106 out of 130 mitotic objects, or (81% predictive, 91% of identified mitotics). Out of 917 single objects, the analysis incorrectly identified only 10 non-mitotics as mitotics (1% total, 8% of identified mitotics); 14 mitotics as interphase (1.4% total, 1% interphase). An automated classification system that would automatically assign values to each object using these or other measurement features can thus be developed, utilizing the principles set forth herein.

In a second experiment, the effects of Taxol on MDCK cells and the different types of morphological effects were observed. A plurality of MDCK cells grown in 96 well plates were treated with Taxol for 4.5 hours at different concentrations (10 uM-1pM). They were then fixed, labeled with Hoechst, and imaged.

This experiment used a labeling protocol comprising: MEOH fix at – 20°, Wash in PBS, Block in PBS/BSA/Serum/Triton-X 100, Incubate with 5 µg/ml Hoechst 10 minutes, and wash.

Cells were inspected for different morphologies and manually counted at each different drug concentration in one well. Fig. 9 shows example images from each drug concentration and the different types of morphologies and cells are highlighted. Fig. 10 shows the distribution of each morphology within the cell population as a function of drug concentration. The higher the concentration of Taxol, the larger proportion of cells underwent apoptosis, and the fewer number of normal mitotic cells were detected.

In a third experiment, the purpose was to determine whether the automated analysis methods developed in the first experiment can detect differences in Hoechst morphology in the presence of 6 known compounds at one concentration and exposure time in one cell line. In this experiment, HeLa cells were separately treated with 6 compounds with known mechanism of action. The quantitative methods described in the first experiment were applied to the Hoechst images.

Approximately 5,000 HeLa cells per well were plated in a Costar black-walled 96 well tissue culture treated plate and left to recover in the incubator for 24 hours. After this time, 10 ug/mL of cytochalasin D (CD), Taxol, hydroxyurea, vinblastine, nocodazole, and staurosporine was added to different wells at a 1:100 addition in DMSO.

The cells were incubated in the presence of drug for 24 more hours. After 24 hours, the cells were removed and fixed as in the first experiment. Then, 9 images per well were collected of the Hoechst staining using a 10x objective.

5 The low magnification images taken of Hoechst were run through the automated image analysis method described in the first experiment. Plots of the average intensity and area were made of each compound. Fig. 11 shows the scatter plots of the compounds. The scatter plots of each compound are visually distinct. For example, cells treated with CD are smaller than control, and cells treated with Hydroxyurea are larger and brighter. Furthermore, the number of cells per well was
10 very different (data not shown).

The effects of different compounds can be clearly and automatically distinguished by identifying changes in cellular morphology. This method can also be used to count adherent cells.

The next experiment was to develop clustering algorithms that assign
15 statistically meaningful values to the representative two dimensional data shown in Fig. 10, and even more complicated clustering of all of the multidimensional data that can be extracted across one, and multiple images.

A fourth experiment was performed to obtain high magnification images of two markers in the presence of drugs. In this experiment, HeLa cells were
20 treated with 80 generic compounds with known mechanism of action. The quantitative methods described in the first experiment were applied to the Hoechst images.

Approximately 5,000 HeLa cells per well were plated in a Costar black walled 96 well tissue culture-treated plate and left to recover in the incubator for 24
25 hours. After this time, 10 ug/mL of each compound from the Killer Plate from Microsource Discovery Systems (Gaylordsville, CT) was added to different wells at a 1:100 addition in DMSO. The cells were incubated in the presence of drug for 24 more hours. After 24 hours, the cells were removed and fixed as in the first experiment. In addition to being labeled with Hoechst 33342 (against chromatin),
30 cells were also labeled with 1 unit of rhodamine-conjugated phalloidin (against actin) for 30 minutes.

The 96 well plate was imaged twice. Once, 9 images per well were collected of the Hoechst staining using a 10x objective. After this, one image per well of both the phalloidin and Hoechst staining was collected using a 40x objective.

The resulting high magnification images were analyzed qualitatively and distinct pattern differences were detected in both the Hoechst and phalloidin images. Fig. 12 shows three example images from the experiment. The top row is the Hoechst staining, and the bottom row is the phalloidin staining from the same well. The columns show the images from wells treated with just DMSO (control), cytochalasin D, and Colchicine. The morphology of each marker is different in the presence of each drug. Interestingly, there is an effect in the morphology of the chromatin in the Hoechst image of cytochalasin D, which directly targets the actin cytoskeleton (and thus there is an expected effect in the phalloidin image). Also, there is an effect on the actin cytoskeleton, compared to control, in the presence of colchicine that directly targets the microtubule network.

The low magnification images were analyzed as described in the first experiment, and different patterns were seen in both the average intensity vs. area plots, and in the number of cells per well (data not shown). Thus, changes in patterns of a marker that is "down-stream" from the direct target of a compound are detectable. Automated image analysis protocols for actin and other markers can be developed similarly, again utilizing the principles set forth herein.

A fifth experiment was performed to test quadruple labeling of 9 different cell lines grown in normal conditions. In this experiment, NCI-H460, A549, MDA-MD-231, MCF-7, SK-OV-3, OVCAR-3, A498, U-2 OS, and HeLa cells were plated. Then, the cells were fixed and stained for portions of the each cell known as DNA, tubulin, actin, and Golgi.

The following table summarizes the procedures for this experiment:

Action	Active Ingredient/Notes	Buffer	Vol/ well	Desired Time	Temp
Remove media	NOTE: gently by pipetting, not aspiration				
Fix	4% Formaldehyde	PBS	100µl	20 min	rt
Wash		TBS	100µl	5 min	rt
Wash		TBS	100µl	5 min	rt

Permeabliz e	0.1% Triton X-100	TBS	100μl	10 min	rt
Permeabliz e	0.1% Triton X-100	TBS	100μl	10 min	rt
Block	% BSA % Serum Filter sterilize before use	TBS w/azide	100μl	1hr or o/n	rt or 4°C
Primary Antibody	1:1000 dilution of DM1α	TBS + 1% BSA + 0.1% TX-100	50μl	1hr or o/n	rt or 4°C
Wash		TBS	100μl	5 min	rt
Wash		TBS	100μl	5 min	rt
Wash		TBS	100μl	5 min	rt
Fluorescent Stain	FITC lens culinaris 1:500 Rhodamine-Phalloidin 1:500 CY5 goat anti-mouse 1:100	TBS + 1% BSA + 0.1% TX-100	50μl	1 hr.	rt, dark
Wash		PBS	100μl	5 min	rt, dark
Hoechst	1:1000 dilution of 5mg/ml	TBS	100μl	15 min	rt, dark
Wash		PBS	100μl	5 min	rt, dark
Wash		PBS	100μl	5 min	rt, dark
Wash		PBS	100μl	5 min	rt, dark
Store		PBS	200μl	1 month	4°C

Cells were plated out at different densities for 48 hours. Cells were fixed and labeled by the above method. Cells were imaged using an automated imaging system that collected 9 images from each marker using a 10x objective.

Higher magnification images were collected of a few cells for demonstration purposes.

In this experiment, each cell line demonstrated different morphological patterns as determined by phase. For example, A549 cells are much more compacted than OVCAR-3 cells as determined by phase contrast imaging (data not shown). The different fluorescent markers showed even bigger differences between different cell lines. Figs. 13 and 14 show 4 panels of each marker for A549 (Fig. 13) and OVCAR-3 cells (Fig. 14). The markers are Hoechst (upper left), Phalloidin (upper right), Lens culinaris (lower left), and DM1a antibody (lower right). The following table summarizes the qualitative differences between these images:

MARKER	A549	OVCAR3
Hoechst/DNA	small	large
Phalloidin/actin	fuzzy	crisp - many stress fibers
Lens culinaris/Golgi	compact	Disperse/punctate
DM1alpha/Tubulin	perinuclear	evenly distributed

Higher magnification images were taken of the OVCAR3 cells. Fig. 15 shows the same markers at 20x, and Fig. 16 shows the markers at 40x. While the highest magnification images show the most detail, these images illustrate that very little morphological or feature information is lost in the 10x images.

These data exemplify the differences in morphology seen between different cell types. Thus the automated image analysis software can be customized for each marker in each cell type. Different drugs should effect these morphologies differentially.

An automated quantification method for each marker and cell line can be similarly developed.

A sixth experiment was conducted with a more sophisticated software package and to develop more flexible image recognition algorithms. In this experiment, prototype image features extraction was performed using MatLab programming language with image toolbox and SDC morphology toolboxes. Algorithms are being developed that will automatically identify objects on images and

to measure various morphological and feature parameters of these objects. Many different features for each of the cellular markers were acquired.

An example of a MatLab program called "AnalyseDNA" that takes as an input an unlimited number of images, identifies individual objects in these images
 5 based on either their intensities, or based on edge-detection algorithms, and extracts a number of morphological and intensity characteristics of these objects. A copy of this program follows:

Listing of the AnalyseDNA.m program and of some of the supporting subroutines

10

```
function files_analysed = AnalyseDNA(filemask, outpath,
nx, ny, filter_range, dext, modifier, sfname)
% AnalyseDNA performs measurements on files of DNA images
% V1. EV 2-11-99; 2-15-99; 2-16-99
```

15

```
%
% files_analysed = AnalyseDNA(filemask, outpath, nx, ny,
filter_range, dext, modifier, sfname)
%
% PARAMETERS:
```

20

```
% ALL PARAMETERS ARE OPTIONAL
```

```
%
```

```
% FILEMASK - mask for file names to be analyzed
INCLUDING PATH(for example c:\images\*.tif)
```

25

```
% DEFAULT '.*.tif' (all *.tif files in the current
directory).
```

```
%
```

```
% OUTPATH - path to a directory where all the output
files will be placed.
```

```
% DEFAULT - output is saved in the same directory
```

30

```
which contains images
```

```
%
```

```
% NX, NY - number of individual images in montage
images along X and Y axes (DEFAULT 1)
```

```
%
```

```
%    FILTER_RANGE - 3 col-wide array (or[]). Specifies
how data is filtered when summary is calculated
%    this parameter internally is passed to GetDNADData
and then to GetSummaryData - see these
5 %    functions for details. For example: [2 2 Inf; 6 100
8000] will case all raws of data for which
%    values in column 2 are less than 2 and all raws
where values in column 6 are less than 100 or
%    more than 8000 to be excluded from all
10 calculations of a summary.
%    DEFAULT - [] (means do not filter, summarize all
data)
%
%    DEXT - string. Extension for data files being saved.
15 %    DEFAULT 'dat';
%
%    MODIFIER - this modifier is 'SUMMARY', summary file
is created;
%    'SUMMARY ONLY' - only summary is generated,
20 data for individual files are not saved
%
%    sfname - string. File name of a summary file
%    DEFAULT 'summary[date].dat'
%
25 % OUTPUT:
%
%    AnalysedDNA works on image files or montages. For
each image file it creates a tab-delimits file of
measured
30 %    parameters of all the objects in the montage with
the same base name as a montage file and extension
specified
```



```
%    by dext parameter (or .dat by default) and file
'errors[date].err' - with the list of files that matched
the
%    filemask but could not be processed.
5 %    If 'summary' or 'summary only' modifier is
specified, it also creates a single file
'summary[date].dat' (or
%    different extension, if specified by DEXT) which
contains summary information for all analyzed files.
10 %
%    ALL OUTPUT FILES are saved in a directory specified
by OUTPATH parameter
%
%    RETURNS *files_analysed* - number of files that have
15 been successfully processed.
%
%    Column designations in the output files are
described in GetDNADData
%
20 % FILE NAME CONVENTIONS
%    AnalyseDNA attempts to identify a number for each
file to identify the file in summary output.
%    It does that by looking for the first space or
underscore, followed by a number and then takes
25 %    as many successive numbers as it can find. If it
fails to identify a number it assigns a
%    default which is -1
%
%
30 % SEE ALSO GetDNADData, GetSummaryData
%
% TO DO    improve error handling in opening and writing
files (GLOBAL error_file ?)
```

```
%          include procedures for writing text headers
into the output files

if nargin > 8
5   error ('Wrong number of input parameters');
end
if nargsout >1
    error ('Wrong number of output parameters: only one
allowed');
10 end

% set defaults
need_summary = 0;
summary_only = 0;
15 use_default_outpath = 0;
datestring = datestr(floor(now));
if nargin == 7      % set default summary file name
    sfname = ['summary' deblank(datestring)]; % extension
will be appended later based on dext
20   if deblank(upper(modifier)) == 'SUMMARY'
        need_summary = 1;
    elseif deblank(upper(modifier)) == 'SUMMARY ONLY'
        need_summary = 1;
        summary_only = 1;
25   else
        error (['Wrong parameter: unknown modifier '
modifier]);
    end
end
30
if nargin == 5
    % default data file extension
    set dext = 'dat';
end
```

```
    if nargin == 4
        % default filter range
        filter_range = [];
    end
5   if nargin == 3
        ny = 1; % default number of images in montage along Y
    end
    if nargin == 2
        nx = 1;
10   end
    if nargin == 1
        use_default_outpath = 1;
    end
    if nargin == 0
15     filemask = '*.tif'
    end

    % check parameters
    if ( ~ischar(filemask) | ~ischar(dext) | ~ischar(sfname)
20   )
        error('Wrong parameter type: filename, filepath,
dext and sfname should be strings');
    end
    if ( ( size(nx) ~= [1 1] ) | ( size(ny) ~= [1 1] ) )
25     error ('Wrong parameter type: nx and ny should be
scalars (1x1 arrays)');
    end
    if (~isempty(filter_range) & size(filter_range, 2) ~= 3)
        error ('Wrong parameter type: filter range should be
30  [] or 3 - cols-wide array');
    end
    % end testing parameters

    % Generate list of files to process
```

```
datapath = getpath(filemask);
if use_default_outpath == 1
    outpath = datapath;
5 end
if exist(outpath, 'dir') ~= 7
    error(['Path ' outpath, 'not found. Exiting..']);
elseif exist(datapath, 'dir') ~= 7
    error(['Path ' datapath, 'not found. Exiting..']);
10 end

sfname = makefullname(outpath, sfname, dext);
if need_summary == 1
    if exist(sfname, 'file')
15     disp(['File ', sfname, 'already exists!']);
    input ('Press ^C to abort, Enter to delete and
continue');
    delete(sfname);
    end
20 end

flist = FileList(getfname(filemask), datapath);
numfiles = size(flist, 1); % total number of files to
25 process
disp(['About to process ', num2str(numfiles), ' files']);
%DEBUG - commented out "input" to run from Wrod
input('Press ^C to abort, Enter to continue');

30 % main loop where the job gets done:
error_file = makefullname(outpath, ['error' datestring
'.err']);
num_processed = 0;
num_error = 0;
```

```
for i = 1:numfiles
    % first generate file name for a data output file
    current_fullname = flist(i, :); % full name with path
    and extension
5    current_datafile = makefullname(outpath,
    makefname(getbasefname(current_fullname), dext) );

    %extract number from a filename
    fnumber = getfilenumber(current_fullname);
10

    % load an imagefile, record errors
    read_error = 0;
    try
        I = imread(current_fullname);
15        %DEBUG
        disp(['Image file #', num2str(fnumber), '
loaded']);
    catch
        % record file-opening error in an error_file
20        read_error = 1;
        num_error = num_error +1;
        msg = [current_fullname ': ' lasterr];
        add_error_msg(error_file, msg);
    end

25
    % extract and write data to a file in outpath
    if read_error ~=1
        if (need_summary == 0)
            %DEBUG
30            disp(['Starting analysis of file #',
num2str(fnumber), '.']);
            current_data = GetDNADData(I, nx, ny, fnumber);
            %DEBUG
```

```

        disp (['Finished analysis of file #',
num2str(fnumber), '.']);
        %load current_data.mat 'current_data';
        write_data(current_data, current_datafile);
5      else      %summary needed
        %DEBUG
        [current_data, current_summary] = GetDNADData(I,
nx, ny, fnumber, filter_range);
        %load current_data.mat 'current_data';
10      %load current_summary.mat 'current_summary';
        write_summary (current_summary, sfname);
        if summary_only ~= 1
            write_data(current_data, current_datafile);
        end
15      end
    end
end % of the main for loop
num_processed = numfiles - num_error;

20 %=====end function AnalyseDNA()
=====

%=====
=====

25 function result = add_error_msg(filename, msg)
% adds string MSG to an errorfile FILENAME
% returns 1 if success, 0 if failure

err_FID = fopen(filename, 'at');
30 if err_FID == -1
    warning(['Can not open error file ' filename]);
else
    fprintf(err_FID, '%s\n', msg);
    fclose(err_FID);

```

```

end
%=====end function add_error_masg()
=====

5 %=====
=====

function N = getfilenumber(fname)
% returns the first number extracted from a file name
(string) or -1 if fails to extract any number
10 numbers = NumbersFromString( getfname(fname) ); % vector
of all numbers encoded in the name

                                % (but not in the path, even if
present)
15 if isempty(numbers)
    N = (-1);    % return -1 if no numbers found in the
name
else
    N = numbers(1);
20 end

%===== end function getfilenumber()
=====

25 %=====
=====

function result = write_data(data_array, file_name)
% writes data in a data_array in a tab-delimited ascii
file.
30 % result is 0 if success and -1 if failure
% if file_name exists, overwrites it
result = -1;
try
    fid = fopen(file_name, 'wt');

```

```

        if fid ~= -1
            for k = 1:size(data_array, 1)
                fprintf(fid, '%g\t', data_array(k, :));
                fprintf (fid, '\n');
5         end
            test = fclose(fid);
            result = -1;
        catch
            result = -1;
10    end

%===== end function write_data()
=====

15 %=====
=====
function result = write_summary (s_vector, file_name)
% appends summary vector s_vector to a file_name (ASCII
tab-delimited file).
20 % if file_name does not exist, creates it.
% result is 0 if success and -1 if failure
%
result = -1;
try
25     % debug
        fid = fopen(file_name, 'at');
        result = fprintf(fid, '%g\t', s_vector);
        result = fprintf(fid, '\n');
        result = fclose(fid);
30     result = 0;
    catch
        result = -1;
    end
end

```



```

% ===== end function write_summary()
=====

function Data = GetObjectsData(I, Ilabel)
5 % GetObjectsData returns array measurements of objects in
  image "I" masked by "Ilabel"
  % EV 2-3-99; 2-10-99
  % OData = GetObjectsData(I, Ilabel) returns an array of
  morphological and intensity measurements
10 %   taken from a grayscale image "I". Objects are
  identified on a mask image Ilabel, usually
  %   created by bwlabel()
  % OUTPUT:
  % Each row in the output array OData represents
15 individual object
  % columns contain the following measurements:
  %
  %   1 - Index ("number" of an object);      8 -
  Solidity;
20 %   2 - X coordinate of the center of mass; 9 - Extent;
  %   3 - Y coordinate      "-"; 10 - Total
  Intensity;
  %   4 - Total Area (in pixels);              11 - Avg.
  Intensity;
25 %   5 - Ratio of MajorAxis/MinorAxis;        12 - Median
  Intensity;
  %   6 - Eccentricity;                        13 - Intensity of
  20% bright pixel
  %   7 - EquivDiameter;                       14 - Intensity of
30 80% bright pixel
  %
  % For details on morphological parameters see information
  on MatLab imfeature();

```

```
% Intensity parameters are either obvious or are
documented in comments in this file.

% Procedures in this file are documented in notebook file
"MATLAB Measuring Nuclei (1) 1-29-98.doc"

5
if (nargin ~= 2)
    error ('function requires exactly 2 parameters');
end
if (nargout ~= 1)
10    error ('function has 1 output argument (array X by
    14)');
end

% finished checking arguments

15
% first collect morphological parameters in a structure
array:
ImStats = imfeature(Ilabel, 'Area', 'Centroid',
    'MajorAxisLength',...
20    'MinorAxisLength', 'Eccentricity', 'EquivDiameter',
    ...
    'Solidity', 'Extent', 8 );

% now convert it into array (matrix) while collecting
25 intensity data for each object:

%preallocate output array:
numobjects = size(ImStats, 1);
OData = zeros(numobjects, 14);
30 %now convert ImStats into array and add intensity data to
it
for k=1:numobjects
    OData(k, 1) = k;
    OData(k, 2) = ImStats(k).Centroid(1);
```

```

        OData(k, 3) = ImStats(k).Centroid(2);
        OData(k, 4) = ImStats(k).Area;
        OData(k, 5) = (ImStats(k).MajorAxisLength) /
        (ImStats(k).MinorAxisLength);
5         OData(k, 6) = ImStats(k).Eccentricity ;
        OData(k, 7) = ImStats(k).EquivDiameter;
        OData(k, 8) = ImStats(k).Solidity;
        OData(k, 9) = ImStats(k).Extent;

10         % now collect and assign intensity parameters from
        image I

        object_pixels = find( Ilabel == k);
        object_area = size(object_pixels, 1); %same as total
15 number of pixels in the object
        object_intensities = double(I(object_pixels)); %
        need to convert to double to do math
        sorted_intensities = sort(object_intensities); %
        will need to get median, 20% and 80% pixels
20         total_intensity = sum(object_intensities, 1);
        avg_intensity = total_intensity / object_area;
        median_intensity = sorted_intensities( floor(
        object_area/2 ) + 1 );
        pix20 = sorted_intensities( floor(object_area*0.2)+1
25 ) ; %brightest pixel among dimmest 20%
        pix80 = sorted_intensities( floor(object_area*0.8)+1
        ) ;

        OData(k, 10) = total_intensity;
30         OData(k, 11) = avg_intensity;
        OData(k, 12) = median_intensity;
        OData(k, 13) = pix20; %brightest pixel among dimmest
        20%

```

```

        OData(k, 14) = pix80; %dimnest pixel among brightest
    20%
    end %for

5   %===== end function
    GetObjectsData()=====

function Imask = MaskDNA1(I);
10  % MaskDNA1 - generates binary mask for cell nuclei
    through edge detection
    % EV 1-22-99; 2-6-99; 2-10-99
    % Imask = MaskDNA1(I)
    % PARAMETERS
15  %   I - intensity image (grayscale)
    % OUTPUT
    %   Imask - BW image with objects from I
    %
    % For more details see Notebook Matlab_DNA_masking1_1-22-
20  99.doc
    % Uses SDC Morphology Toolbox V0.7

    if (nargin ~= 1)
        error('Wrong number of input parameters');
25  end
    if (nargout ~= 1)
        error('Wrong number of output parameters: one output
        argument should be provided');
    end
30

    Imask = edge(I, 'canny');
    Imask = mm dil(Imask, mmsecross(1));
    Imask = mmero ( mmc lohole(Imask,mmsecross(1)));

```

```

Imask = mmedgeoff(Imask, mmsecross(1));
% note that mmedgeoff this command removed FILLED OBJECTS
but not touching OUTLINES.
% these outlines can be removed by filtering:
5  Imask = medfilt2(Imask, [5 5]);

%=====end MaskDNA1 =====

```

Given the list of image files or montages of images as an input, this
 10 program creates an individual file for each image that contains the following
 quantitative measurements for all objects identified in the image:

1 - Index ("number" of an object);	8 - Solidity;
2 - X coordinate of the center of mass;	9 - Extent;
15 3 - Y coordinate "-";	10 - Total Intensity;
4 - Total Area (in pixels);	11 - Avg. Intensity;
5 - Ratio of MajorAxis/MinorAxis;	12 - Median Intensity;
6 - Eccentricity;	13 - Intensity of 20% bright pixel
7 - EquivDiameter;	14 - Intensity of 80% bright pixel

20 A fragment of an output for a single file, containing 9 images of cells
 stained for DNA and acquired with a 10x objective. A montage image that was used
 as a source to generate data in A is presented in Fig. 17.

The same program also summarizes measurements across many files
 and performs statistical analysis of the summary data. It creates a summary file with
 25 the following data:

1 - Image file number;	
2 - Average object Area (in pixels);	3 - STD (standard deviation) of
2;	
30 4 - Avg. of Ratio of MajorAxis/MinorAxis;	5 - STD of 4;
6 - Avg. Eccentricity;	7 - STD of 6;
8 - Avg. EquivDiameter;	9 - STD of 8;
10 - Avg. of Solidity;	11 - STD of 10;

- | | |
|---|----------------|
| 12 - Avg. of Extent; | 13 - STD of 11 |
| 14 - Avg. of objects Total Intensity; | 15 - STD of 14 |
| 16 - Avg. of objects Avg Intensity; | 16 - STD of 15 |
| 18 - Avg. of objects Median intensity; | 19 - STD of 18 |
| 5 20 - Avg. of objects intensity of 20% bright pixel; | 21 - STD of 19 |
| 22 - Avg. of objects intensity of 80% bright pixel; | 23 - STD of 21 |

An example of summary output obtained by running AnalyseDNA against 10 montage files also is shown in Appendix B.

10 A seventh experiment was conducted in order to use sequence analysis algorithms to analyze features of cell images. In this experiment, HeLa cells were treated for 24 hours with several different compounds, and then fixed, and stained with a fluorescent DNA dye. One image of these cells was acquired for each of the treatments and morphometric parameters and features were measured:

15 Resulting measurements were arranged into a string of numbers and reduced to a pseudo- nucleic acid sequence using following rules: At any given position in the sequence a number was substituted by "t" (a code for thymidine) if its value is among highest 25% of the values at the corresponding position in the data set, "g" if it is between 50% and 25%, "c" if it is between 75% and 50%, and "a" if it
20 belongs to lowest 25% of values. Thus one descriptor or sequence was generated per treatment as illustrated in Fig. 18.

Resulting sequences were clustered using an AlignX module commercial software package Vector NTI (<http://informaxinc.com>), which uses a Neighbor Joining algorithm for sequence clustering.

25 The resulting dendrogram is presented in Fig 18. On the dendrogram the closest "leafs" correspond to the closest pseudo-sequences. Interestingly, compounds with similar mechanisms of action cluster together on the dendrogram. Another example of the generation of pseudo-sequences and clustering is shown in Fig. 19.

30 In some embodiments, techniques according to the present invention can provide tools for the later stages of drug development such as clinical trial design and patient management. The properties of known drugs such as clinical trial and patient response information will be used in a similar fashion as the pre-clinical

information to provide predictions about the properties of novel compounds. Because the human cell is the locus of drug action, a database containing drug-cell interactions can be able to provide predictive information for this aspect of drug development.

Although the above has generally described the present invention
5 according to specific systems, the present invention has a much broader range of applicability. In particular, the present invention is not limited to a particular kind of data about a cell, but can be applied to virtually any cellular data where an understanding about the workings of the cell is desired. Thus, in some embodiments, the techniques of the present invention could provide information about many
10 different types or groups of cells, substances, and genetic processes of all kinds. Of course, one of ordinary skill in the art would recognize other variations, modifications, and alternatives.

APPENDIX A

EV Table 1.doc

Example of the output of AnalysedNA.m program
(measurements for a single 3 by 3 montage image)

File	Subimage	object	X coord	Y coord	Area	Area ratio	Eccentricity	Equival	Solidity	Extent	Intensity	Avg. Intensity	Median Intensity	201 pla.	ROI pla.
1	1	1	12.2897	152.635	145	1.17293	0.532621	13.5875	0.923367	0.739796	4605	31.7396	31	23	37
1	1	2	16.352	416.032	125	1.60394	0.182471	12.6157	0.905391	0.78125	4606	36.848	38	30	45
1	1	3	20.1073	73.8079	177	1.09485	0.413785	15.0121	0.917098	0.691406	4769	26.9435	29	22	31
1	1	4	21.6186	402.744	47	1.36215	0.478006	7.39978	0.914594	0.707457	3650	85.114	87	67	105
1	1	5	27.0918	164	96	1.30887	0.443134	11.0558	0.888939	0.671329	4502	46.8958	48	38	54
1	1	6	30.3252	359.531	206	2.31006	0.403309	16.1953	0.927928	0.715278	6380	30.9705	33	24	37
1	1	7	33.6429	167.573	89	1.24884	0.471696	10.6451	0.927083	0.711667	4225	47.4719	50	38	44
1	1	8	35.0411	16.9728	166	1.25176	0.401495	13.6347	0.929296	0.745919	5415	37.089	40	29	44
1	1	9	37.766	366.021	41	1.84062	0.439542	7.33578	0.87037	0.632778	6657	141.851	142	113	171
1	1	10	49.1078	170.004	232	1.90491	0.451127	17.187	0.852941	0.70303	5832	42.3193	45	33	51
1	1	11	56.0769	126.531	221	1.95704	0.455955	16.7746	0.924686	0.683335	7040	31.8552	33	25	31
1	1	12	57.7755	41.932	147	1.33627	0.463301	13.6009	0.907407	0.706771	4745	32.415	34	26	39
1	1	13	51.444	366.851	171	2.27223	0.487553	14.7555	0.872449	0.706412	5318	56.8421	56	43	68
1	1	14	56.4079	282.272	208	1.92782	0.451944	16.1953	0.923767	0.63375	7137	34.6156	37	28	41
1	1	15	57.0648	227.176	108	1.73485	0.413089	11.7265	0.913254	0.701299	4644	43	43	51	
1	1	16	66.1714	333.161	315	1.13194	0.437266	20.0267	0.75	0.528756	13151	48.0984	50	36	42
1	1	17	65.1109	402.411	220	1.70147	0.40906	16.7366	0.920502	0.617059	9809	44.5864	46	35	54
1	1	18	71.8449	402.13	185	1.73678	0.424583	15.3476	0.911111	0.619123	6124	33.1027	35	25	39
1	1	19	71.626	164.854	123	1.71580	0.417622	12.5143	0.911111	0.723229	4811	39.3577	41	30	47
1	1	20	71.4869	132.513	306	1.6379	0.791981	19.7386	0.622231	0.622231	14559	36.7459	40	38	57
1	1	21	76.7377	208.27	122	1.3257	0.467941	12.4634	0.910448	0.735394	4483	40.0513	43	37	43
1	1	22	81.786	52.9612	117	1.44713	0.791616	12.2053	0.846361	0.62857	4686	40.0513	43	37	43
1	1	23	88.7292	281.574	373	2.17388	0.481916	21.7926	0.841891	0.513219	16109	43.1877	46	31	52
1	1	24	88.7165	301.976	85	1.20799	0.450931	10.4031	0.914628	0.708333	5369	53.9882	57	43	63
1	1	25	88.1408	176.231	143	1.43573	0.717545	13.4935	0.940789	0.794444	4878	34.1119	35	27	41
1	1	26	91.4229	376.924	170	1.36852	0.492469	16.7123	0.833333	0.833333	4933	29.0176	30	23	35
1	1	27	97.7604	371.795	288	1.92353	0.462119	19.1492	0.9	0.606316	10663	37.0243	39	29	45
1	1	28	95.5841	330.343	113	1.09825	0.413609	11.9948	0.856825	0.608363	4560	40.177	43	32	48
1	1	29	95.9932	248.602	118	1.2774	0.472219	12.2573	0.921675	0.746234	4873	41.2866	43	32	51
1	1	30	103.37	327.502	124	1.66415	0.718927	12.6574	0.945736	0.813233	6652	38.2212	40	31	47
1	1	31	103.37	327.502	124	1.3208	0.753111	13.0619	0.917808	0.671179	4358	32.5224	34	27	38
1	1	32	103.37	327.502	124	1.30329	0.450449	12.2573	0.907692	0.694118	4695	39.7181	42	30	40
1	1	33	121.22	265.08	328	1.20328	0.405916	20.7734	0.900552	0.624923	15664	48.4626	50	39	60
1	1	34	121.22	265.08	328	1.51045	0.757011	13.9988	0.927632	0.731375	6429	45.5857	49	37	57
1	1	35	127.84	60.3355	132	1.75689	0.422208	11.9116	0.921212	0.767677	6875	45.2303	47	36	54
1	1	36	137.083	128.093	266	1.445	0.440315	16.4033	0.796407	0.538462	9810	36.8197	38	26	45
1	1	37	137.083	128.093	266	1.445	0.440315	16.4033	0.937143	0.788462	7337	44.7378	47	35	53
1	1	38	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	39	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	40	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	41	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	42	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	43	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	44	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	45	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	46	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	47	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	48	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	49	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	50	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	51	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	52	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	53	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	54	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	55	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	56	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	57	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	58	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	59	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	60	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	61	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	62	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	63	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	64	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	65	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	66	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	67	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33
1	1	68	137.083	128.093	266	1.39276	0.459732	15.4204	0.935	0.799145	5227	27.9319	27	22	33

EV Table 1.doc

49	243.509	86.7857	224	1.87991	0.816782	16.888	0.899598	0.635344	8827	39.4053	42	30
50	246.831	322.144	160	1.79661	0.820819	16.273	0.91954	0.71555	5025	31.4053	33	32
51	248.234	413.026	71	1.15364	0.898511	9.90149	0.875	0.7	4352	36.3195	39	35
52	255.945	41.816	163	1.58857	0.777002	9.1052	0.91573	0.679167	4720	28.3511	30	32
53	255.945	398.848	66	1.03149	0.245208	9.167	0.916667	0.410815	4440	47.2127	71	32
54	263.092	375.55	251	1.95991	0.860038	17.8769	0.886926	0.597619	10500	41.0227	33	32
55	263.092	375.55	251	1.95991	0.860038	17.8769	0.886926	0.597619	10500	41.0227	33	32
56	264.397	209.402	131	1.31512	0.795552	14.3175	0.916773	0.484014	5136	31.9004	32	32
57	264.397	209.402	131	1.31512	0.795552	14.3175	0.916773	0.484014	5136	31.9004	32	32
58	266.137	346.328	131	1.58185	0.774829	11.8882	0.909722	0.452292	9633	43.7658	75	32
59	276.221	171.24	204	2.05613	0.873763	16.1165	0.918519	0.596651	70531	34.3637	35	32
60	276.221	285.089	287	1.27833	0.622935	19.1176	0.872321	0.450791	10320	36.5551	35	32
61	276.221	391.32	150	1.10321	0.422329	38.8198	0.920245	0.765208	9202	41.3167	35	32
62	276.221	391.32	150	1.10321	0.422329	38.8198	0.920245	0.765208	9202	41.3167	35	32
63	285.305	154.719	231	1.56301	0.748455	17.1439	0.923913	0.417308	4387	31.4118	35	33
64	285.305	203.688	221	1.75368	0.822092	16.7746	0.931432	0.475139	5380	37.1429	35	33
65	284.739	355.022	46	1.74017	0.818394	7.65304	0.867923	0.730159	4986	44.3846	48	34
66	291.4	319.71	145	1.13164	0.618667	13.5875	0.917722	0.733208	4940	31.049	35	32
67	291.451	442.734	192	2.01531	0.888208	15.6353	0.911176	0.793388	5972	31.1042	32	32
68	293.41	389.276	58	1.25774	0.60651	8.59348	0.920815	0.725	3964	68.3448	70	55
69	299.182	285.162	159	1.39289	0.696112	13.2283	0.928815	0.757143	5103	32.0943	33	36
70	300.14	356.347	150	1.31538	0.649643	13.8198	0.925926	0.78123	5369	35.7933	37	30
71	311.5	260.38	382	1.35137	0.726532	22.054	0.943267	0.598746	16117	42.1911	41	32
72	311.5	260.38	382	1.35137	0.726532	22.054	0.943267	0.598746	16117	42.1911	41	32
73	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
74	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
75	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
76	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
77	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
78	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
79	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
80	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
81	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
82	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
83	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
84	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
85	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
86	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
87	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
88	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
89	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
90	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
91	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
92	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
93	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
94	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
95	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
96	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
97	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
98	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
99	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
100	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
101	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
102	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
103	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
104	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
105	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
106	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
107	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
108	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
109	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
110	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
111	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
112	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
113	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
114	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
115	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
116	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
117	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
118	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
119	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
120	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
121	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
122	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
123	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
124	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
125	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
126	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
127	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
128	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
129	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
130	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
131	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
132	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
133	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
134	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	32
135	318.37	332.894	161	1.35063	0.49465	14.3175	0.936047	0.766667	4966	30.8447	33	

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1	136	485.08	156.469	293	2.80072	0.93405	19.5115	0.89571	0.69991	1.0053	47.0401	50	38	57
2	137	488.732	106.247	190	1.75179	0.821058	15.5516	0.858156	0.666667	0.866	46.2421	50	35	57
3	138	492.318	317.287	155	1.824	0.673195	15.751	0.812011	0.71301	5108	27.7133	29	23	34
4	1	10.098	359.018	102	1.38118	0.673195	11.3651	0.918919	0.713281	5508	54	39	58	68
5	2	11.7415	119.534	176	1.8992	0.850152	11.4099	0.88014	0.516471	5099	34.4871	37	25	44
6	3	16.517	274.314	176	1.49674	0.744054	14.3959	0.945116	0.617059	8466	56.0569	60	45	68
7	4	22.1517	323.427	211	1.20929	0.567256	10.2119	0.845116	0.617059	8466	56.0569	60	45	68
8	5	22.7977	351.041	71	1.49712	0.747208	14.2907	0.805579	0.703323	9339	45.9028	47	35	55
9	6	22.7991	366.47	69	1.65511	0.513637	9.70468	0.805579	0.703323	9339	45.9028	47	35	55
10	7	31.5223	50.4478	297	1.47556	0.723029	5.37062	0.873118	0.61897	3983	57.7216	59	44	70
11	8	31.5223	109.832	155	2.56212	0.920487	13.4161	0.838982	0.52002	11786	31.4788	61	31	48
12	9	44.7125	455.738	233	1.9878	0.864216	17.221	0.869101	0.617222	9369	52.1152	42	31	50
13	10	51.9162	275.366	191	1.67447	0.80209	15.5915	0.836714	0.636667	9954	42.4132	34	27	41
14	11	62.4292	262.297	219	2.22958	0.818965	16.6985	0.947205	0.608233	9759	44.5616	46	36	54
15	12	65.7559	171.244	213	2.19727	0.850135	16.4488	0.946667	0.608233	9759	44.5616	46	36	54
16	13	63.4132	488.702	121	1.49032	0.806279	12.4122	0.927264	0.75625	1271	55.1215	56	44	70
17	14	73.3548	400.7762	155	1.30361	0.64153	11.0482	0.927264	0.75625	1271	55.1215	56	44	70
18	15	75	329.552	183	1.97089	0.861719	15.2444	0.915	0.72619	987	27.1421	28	21	31
19	16	80.0786	183.618	210	1.46688	0.794543	16.3518	0.919204	0.714286	7257	34.5571	37	27	41
20	17	79.6596	291.823	141	1.25467	0.603949	13.3988	0.933775	0.783733	6830	34.2551	36	27	40
21	18	80.1049	310.149	224	1.39545	0.697472	12.5651	0.925723	0.751515	6807	37.1532	38	30	44
22	19	86.0366	103.451	184	1.55708	0.746515	11.4503	0.916501	0.69076	5396	32.2927	34	25	39
23	20	90.3161	139.787	174	1.67033	0.800986	11.8843	0.970535	0.654135	4950	26.951	29	21	32
24	21	85.1818	504.591	22	1.58172	0.774787	5.29257	0.916667	0.733333	1517	68.9545	68	61	106
25	22	84.5814	72.4186	43	1.89078	0.848695	7.39928	0.914694	0.671875	6443	149.937	155	105	186
26	23	95.9012	437.878	188	1.46598	0.731051	15.4716	0.930693	0.706767	6330	23.0319	23	18	28
27	24	103.775	259.464	271	2.25576	0.896369	18.5755	0.73842	0.531333	10162	38.2162	40	30	47
28	25	101.716	351.365	109	1.26499	0.680653	11.7406	0.913023	0.676531	4566	61.6899	44	33	52
29	26	100.971	381.4	105	1.8452	0.804731	11.5474	0.913023	0.676531	4566	61.6899	44	33	52
30	27	102.731	24.3077	182	1.42707	0.724519	15.2227	0.938144	0.722222	5335	29.3172	31	24	33
31	28	104.762	241.381	232	1.52487	0.754476	17.9125	0.918182	0.722222	4831	37.1615	37	29	46
32	29	107.462	297.946	130	1.205	0.58271	12.8455	0.928571	0.722222	4831	35.1197	38	28	42
33	30	117.468	139.313	128	1.2113	0.564315	12.7462	0.911176	0.781905	4503	35.1197	38	28	42
34	31	117.476	270.203	165	1.53776	0.759681	13.5073	0.912281	0.615325	622	44.4923	45	34	53
35	32	119.39	474.808	104	1.58258	0.775048	11.2733	0.855615	0.666667	3953	24.7063	25	19	31
36	33	124.631	666.313	160	2.13828	0.883905	11.2733	0.855615	0.666667	3953	24.7063	25	19	31
37	34	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
38	35	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
39	36	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
40	37	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
41	38	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
42	39	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
43	40	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
44	41	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
45	42	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
46	43	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
47	44	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
48	45	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
49	46	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
50	47	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
51	48	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
52	49	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
53	50	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
54	51	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
55	52	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
56	53	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
57	54	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
58	55	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
59	56	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
60	57	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
61	58	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
62	59	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
63	60	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
64	61	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
65	62	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
66	63	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
67	64	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
68	65	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
69	66	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
70	67	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
71	68	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
72	69	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
73	70	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
74	71	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
75	72	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
76	73	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
77	74	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
78	75	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	34
79	76	125.541	692.842	133	1.14759	0.691128	13.0131	0.93007	0.730769	3821	28.7923	30	23	3

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1	1	3	74	270.921	432.627	177	1.27386	0.624112	15.0121	0.917098	0.713692	97.19	55.0791	35	42	69
2	1	3	75	271.481	472.443	221	1.72596	0.613564	16.7716	0.888297	0.613698	97.22	47.3122	45	33	51
3	1	3	76	271.922	505.875	208	2.32522	0.611714	16.2133	0.908297	0.613698	97.22	47.3122	45	33	51
4	1	3	77	281.51	595.384	488	2.07461	0.617653	16.3287	0.916461	0.510435	131.33	51.0102	32	24	38
5	1	3	78	289.331	190.619	144	1.98664	0.613353	13.3006	0.916461	0.510435	131.33	51.0102	32	24	38
6	1	3	79	289.363	256.084	219	2.49567	0.620034	17.8035	0.917125	0.523109	104.86	42.1134	43	33	51
7	1	3	80	290.085	492.403	189	1.23762	0.589177	15.3126	0.921951	0.75	47.3	25.0952	27	20	30
8	1	3	81	293.258	232.6071	112	1.49137	0.741883	11.9416	0.888889	0.622222	44.62	45.2321	47	24	36
9	1	3	82	296.44	278.133	293	1.23274	0.575125	15.3147	0.931299	0.620728	131.63	45.3108	47	24	36
10	1	3	83	296.947	46.0877	114	1.53891	0.759372	12.0078	0.890653	0.690909	45.69	40.2344	40	31	49
11	1	3	84	291.161	204.621	124	1.51011	0.769323	12.5651	0.923194	0.626667	60.13	48.4919	31	39	58
12	1	3	85	291.161	204.621	124	1.51011	0.769323	12.5651	0.923194	0.626667	60.13	48.4919	31	39	58
13	1	3	86	299	473.056	234	1.32743	0.659331	12.8159	0.920298	0.716667	47.3	36.547	38	28	44
14	1	3	87	299	473.056	234	1.32743	0.659331	12.8159	0.920298	0.716667	47.3	36.547	38	28	44
15	1	3	88	299	473.056	234	1.32743	0.659331	12.8159	0.920298	0.716667	47.3	36.547	38	28	44
16	1	3	89	308.782	142.492	238	1.60716	0.793031	11.9948	0.698823	0.620879	50.52	44.708	47	36	54
17	1	3	90	308.782	142.492	238	1.60716	0.793031	11.9948	0.698823	0.620879	50.52	44.708	47	36	54
18	1	3	91	308.782	142.492	238	1.60716	0.793031	11.9948	0.698823	0.620879	50.52	44.708	47	36	54
19	1	3	92	310.453	41.0155	373	1.12602	0.6565	12.7442	0.927536	0.790123	46.66	35.4094	38	28	44
20	1	3	93	310.453	41.0155	373	1.12602	0.6565	12.7442	0.927536	0.790123	46.66	35.4094	38	28	44
21	1	3	94	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
22	1	3	95	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
23	1	3	96	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
24	1	3	97	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
25	1	3	98	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
26	1	3	99	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
27	1	3	100	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
28	1	3	101	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
29	1	3	102	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
30	1	3	103	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
31	1	3	104	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
32	1	3	105	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
33	1	3	106	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
34	1	3	107	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
35	1	3	108	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
36	1	3	109	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
37	1	3	110	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
38	1	3	111	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
39	1	3	112	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
40	1	3	113	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
41	1	3	114	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
42	1	3	115	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
43	1	3	116	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
44	1	3	117	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
45	1	3	118	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
46	1	3	119	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
47	1	3	120	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
48	1	3	121	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
49	1	3	122	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
50	1	3	123	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
51	1	3	124	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
52	1	3	125	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
53	1	3	126	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
54	1	3	127	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
55	1	3	128	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
56	1	3	129	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
57	1	3	130	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
58	1	3	131	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
59	1	3	132	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
60	1	3	133	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
61	1	3	134	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
62	1	3	135	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
63	1	3	136	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
64	1	3	137	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
65	1	3	138	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
66	1	3	139	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
67	1	3	140	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
68	1	3	141	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
69	1	3	142	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
70	1	3	143	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
71	1	3	144	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
72	1	3	145	317.211	458.408	473	1.49285	0.796166	20.2795	0.828205	0.621154	96.95	34.1966	35	25	42
73	1	3	146	317.211</												

EV Table 1.doc

1	1	491.586	286.016	35	1.14928	0.341385	6.67558	0.345916	0.433333	3545	101.284	105	99	117
2	1	491.438	76.3765	130	2.02206	0.471881	12.4655	0.302778	0.433333	4543	35.0231	36	26	43
3	1	499.436	144.308	128	1.24035	0.34184	12.1462	0.320813	0.711111	4856	37.9375	39	30	46
4	1	504.436	285.462	39	1.4304	0.715017	7.04573	0.306971	0.8125	3680	91.359	36	19	113
5	1	22.2001	122.471	152	1.3561	0.475419	12.9116	0.321212	0.710169	4832	31.7895	33	35	39
6	2	22.1391	72.3198	197	1.35473	0.462327	12.4376	0.329245	0.729653	9190	48.1726	30	37	59
7	2	22.1391	140.043	115	1.35777	0.474433	12.1003	0.321933	0.405842	4012	39.2261	47	20	48
8	3	23.2174	287.022	137	1.68419	0.800492	13.1838	0.308639	0.494071	7260	44.1032	40	38	53
9	3	26.6026	344.243	151	1.68495	0.804493	13.1838	0.32378	0.764633	4532	48.0795	50	36	57
10	4	26.0394	162.359	102	1.28299	0.624953	11.3561	0.305941	0.371873	9531	44.1316	45	35	53
11	4	36.734	487.684	182	2.28724	0.906008	15.2444	0.321351	0.371873	9531	34.7182	35	26	43
12	5	39.2551	188.77	173	2.36982	0.939086	21.9052	0.321351	0.419333	13413	26.0928	27	19	31
13	5	49.5199	217.316	177	2.90987	0.731315	12.3092	0.329348	0.535121	10090	40.9866	35	26	40
14	6	40.3109	306.084	119	1.46633	0.731315	12.3092	0.329348	0.535121	10090	38.4739	41	39	49
15	10	49.8124	124.27	259	2.13833	0.803911	14.1555	0.329348	0.535121	10090	28.2573	51	20	40
16	12	53.6559	76.3110	166	1.22662	0.379108	15.389	0.364162	0.761513	9169	41.0021	42	29	51
17	13	61.75	496.487	156	1.3807	0.695518	14.0925	0.324571	0.714706	4601	48.475	38	34	56
18	14	61.8138	110.813	96	1.36163	0.765055	11.0359	0.324571	0.714706	4601	33.9419	35	27	41
19	15	70.4153	190.465	172	2.30563	0.901007	14.7286	0.300526	0.532508	5938	51.7037	38	43	55
20	16	69.6514	97.7718	81	1.41608	0.711712	10.1354	0.3	0.75	4131	42.7843	46	33	52
21	17	87.8529	100.745	102	1.86101	0.843364	11.3961	0.394737	0.706333	4364	40.4701	43	32	46
22	18	89.9759	68.9148	100	1.14834	0.491595	11.7265	0.315234	0.755245	4397	40.7158	42	32	46
23	19	98.4146	183.474	190	2.31245	0.901662	15.5536	0.32233	0.68806	7736	42.9761	46	34	51
24	20	98.4146	173.219	251	1.39113	0.655175	17.8769	0.333086	0.747028	10787	42.2162	44	33	51
25	21	96.8018	253.413	111	1.36667	0.605287	11.8882	0.317355	0.74	4686	42.9752	41	29	46
26	22	96.6612	405.521	221	1.31238	0.648287	12.4122	0.316667	0.765152	4595	42.9752	41	29	46
27	23	98.7228	41.5214	101	1.22463	0.576493	11.3101	0.30991	0.765152	4595	42.9752	41	29	46
28	24	100.185	300.07	115	1.48194	0.718011	14.9696	0.326316	0.735996	5059	38.4443	29	22	34
29	25	101.765	100.185	176	1.86325	0.813776	12.1005	0.326316	0.735996	5059	38.4443	29	22	34
30	26	108.123	141.793	135	1.04071	0.333452	13.1106	0.318347	0.741358	4866	38.0146	36	28	45
31	27	118.773	642.208	18	2.78411	0.895059	7.41764	0.318347	0.741358	4866	38.0146	36	28	45
32	28	124.705	39.2679	112	1.93296	0.83896	11.9416	0.318347	0.741358	4866	38.0146	36	28	45
33	29	132.516	339.373	110	1.56515	0.769275	11.8745	0.32437	0.705128	4808	40.8	43	33	49
34	30	130.016	410.445	128	1.20716	0.560153	12.7662	0.307801	0.761905	4834	38.6719	37	29	45
35	31	130.461	60.240	125	1.23859	0.676922	12.6157	0.325928	0.751576	4636	37.088	38	28	45
36	32	135.846	109.582	182	1.13873	0.531436	15.2229	0.324571	0.764706	5123	28.1444	29	22	34
37	33	140.516	124.595	164	1.60308	0.801354	14.4503	0.321318	0.759259	4936	30.2195	31	24	36
38	34	141.5	164.779	104	1.25544	0.606328	11.5073	0.312281	0.722222	4104	42.3462	44	33	50
39	35	151.376	245.402	128	1.35544	0.675056	12.7662	0.307801	0.711111	4698	38.0146	36	28	45
40	36	154.318	472.929	157	1.45981	0.788134	14.1386	0.326994	0.644444	4545	39.181	38	29	49
41	37	157.259	431.776	116	1.53578	0.758963	13.3512	0.315033	0.729167	4828	38.4837	35	27	47
42	38	157.419	292.534	160	1.03316	0.232051	14.0935	0.321077	0.712657	9109	61.5962	64	49	73
43	39	160.554	163.442	156	1.71923	0.813434	12.4122	0.326296	0.664835	4553	37.8281	39	29	46
44	40	163.455	141	121	1.72672	0.815232	8.95623	0.313043	0.63	7016	111.365	116	91	136
45	41	164.635	45.5079	43	1.63955	0.792517	10.2179	0.313043	0.63	7016	111.365	116	91	136
46	42	168.561	328.11	82	1.63955	0.792517	10.2179	0.313043	0.63	7016	111.365	116	91	136
47	43	171.548	93.6129	341	1.57991	0.721772	20.8369	0.323741	0.645833	16949	43.8387	45	33	55
48	44	170.747	210.378	61	1.21347	0.56884	10.5348	0.315789	0.750909	4778	54.9195	56	43	65
49	45	172.538	132.164	170	1.4118	0.72038	12.8635	0.315493	0.714286	4816	37.0462	37	29	46
50	46	172.538	132.164	170	1.03532	0.256937	10.1554	0.31166	0.81	4816	59.4568	60	45	71
51	47	187.788	395.157	194	1.03532	0.256937	10.1554	0.31166	0.81	4816	59.4568	60	45	71
52	48	187.788	395.157	194	1.03532	0.256937	10.1554	0.31166	0.81	4816	59.4568	60	45	71
53	49	201.162	225.324	579	1.57599	0.722905	17.1515	0.311748	0.714648	5054	26.0515	27	20	32
54	50	201.162	225.324	579	1.57599	0.722905	17.1515	0.311748	0.714648	5054	26.0515	27	20	32
55	51	208.346	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
56	52	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
57	53	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
58	54	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
59	55	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
60	56	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
61	57	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
62	58	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
63	59	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
64	60	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
65	61	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
66	62	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
67	63	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
68	64	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
69	65	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
70	66	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
71	67	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
72	68	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
73	69	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
74	70	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
75	71	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53
76	72	214.391	15.4718	142	1.38993	0.69916	14.0028	0.333333	0.802083	7416	41.4591	42	31	53

EV Table 1.doc

1	308.012	29.431	170	1.2847	0.62778	14.7123	0.202091	0.574603	4598	27.0171	28	21	33
2	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
3	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
4	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
5	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
6	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
7	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
8	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
9	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
10	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
11	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
12	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
13	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
14	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
15	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
16	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
17	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
18	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
19	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
20	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
21	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
22	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
23	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
24	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
25	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
26	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
27	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
28	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
29	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
30	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
31	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
32	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
33	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
34	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43
35	305.881	27.524	126	1.29041	0.61616	12.566	0.1931617	0.5	4480	35.5556	38	28	43

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1	1	171.584	470.016	750	1.71834	0.813219	17.8412	0.919168	0.657895	5098	20.392	21	16	25
2	2	174.47	174.326	115	1.36792	0.602339	12.1005	0.691473	0.654597	4161	36.1874	36	28	35
3	3	177.392	331.348	153	1.70893	0.810918	13.9573	0.910714	0.708333	4100	28.7582	29	22	44
4	4	179.828	341.081	99	1.51368	0.750701	11.2372	0.805684	0.640523	4248	47.1148	49	37	59
5	5	182.462	374.305	196	1.64808	0.804616	15.7973	0.808574	0.670571	3372	23.2281	24	18	24
6	6	184.981	415.42	171	1.60258	0.78143	12.7355	0.814728	0.670455	4074	34.1017	36	26	42
7	7	187.381	415.42	118	1.64552	0.844191	12.2573	0.923423	0.716783	4596	22.4195	24	17	28
8	8	189.828	415.42	118	1.64552	0.813525	16.1558	0.89916	0.629412	4016	37.5327	38	28	41
9	9	192.333	40.8098	205	2.43432	0.913525	16.1558	0.89916	0.629412	4016	46.7293	47	37	55
10	10	194.827	497.308	107	1.31808	0.689916	15.1808	0.918782	0.709504	8458	15.0994	47	37	55
11	11	197.308	246.271	181	1.20958	0.562581	15.1808	0.918782	0.709504	8458	44.7293	47	37	55
12	12	200.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
13	13	203.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
14	14	205.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
15	15	208.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
16	16	210.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
17	17	213.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
18	18	215.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
19	19	218.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
20	20	220.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
21	21	223.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
22	22	225.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
23	23	228.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
24	24	230.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
25	25	233.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
26	26	235.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
27	27	238.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
28	28	240.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
29	29	243.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
30	30	245.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
31	31	248.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
32	32	250.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
33	33	253.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
34	34	255.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
35	35	258.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
36	36	260.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
37	37	263.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
38	38	265.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
39	39	268.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
40	40	270.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
41	41	273.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
42	42	275.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
43	43	278.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
44	44	280.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
45	45	283.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
46	46	285.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
47	47	288.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
48	48	290.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
49	49	293.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
50	50	295.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
51	51	298.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
52	52	300.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
53	53	303.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
54	54	305.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
55	55	308.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
56	56	310.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
57	57	313.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
58	58	315.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
59	59	318.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
60	60	320.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
61	61	323.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
62	62	325.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
63	63	328.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
64	64	330.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
65	65	333.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
66	66	335.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
67	67	338.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
68	68	340.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
69	69	343.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
70	70	345.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
71	71	348.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
72	72	350.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
73	73	353.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
74	74	355.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
75	75	358.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
76	76	360.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
77	77	363.333	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
78	78	365.827	310.168	105	1.63706	0.781746	14.6415	0.918782	0.709504	8458	44.7293	47	37	55
79	79	368.333	310.168	105	1.63706	0.781746	14.6415	0.918782</						

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106	359.018	312.36	114	1.70064	0.872412	32.0078	0.597638	0.574374	41.91	36.6754	38	28	43
107	361.716	169.031	129	1.60135	0.791018	32.8159	0.508428	0.461538	75.36	59.4166	60	46	71
108	370.42	126.786	132	1.12766	0.42473	31.9018	0.503226	0.711948	42.18	37.4607	30	30	64
109	379.866	101.258	91	1.21577	0.568727	31.3132	0.503226	0.711948	41.16	42.7276	44	34	56
110	382.727	601.684	187	1.718828	0.820359	35.4306	0.921215	0.611111	68.30	46.1697	48	36	57
111	388.96	193.188	303	1.87034	0.830533	39.6616	0.921215	0.611111	68.30	36.9967	38	29	65
112	397.155	337.23	174	1.72598	0.812333	33.1559	0.913789	0.43125	87.51	50.3276	52	39	61
113	384.648	261.441	136	1.69281	0.804233	33.1559	0.913789	0.43125	87.51	33.6566	35	27	61
114	401.428	302.432	144	1.51937	0.751059	22.9591	0.866109	0.613333	136.64	47.3459	50	32	61
115	395.517	232.544	149	2.01919	0.872845	33.7736	0.923666	0.613333	51.11	36.3154	38	29	44
116	414.597	773.038	384	1.50902	0.748301	22.3116	0.829376	0.613333	199.55	28.0288	21	20	32
117	424.436	70.7462	159	1.6214	0.790105	34.2789	0.910828	0.613333	45.92	28.8805	29	22	35
118	429	16	109	1.24326	0.581659	31.7806	0.921071	0.712333	39.03	35.8033	37	30	42
119	431.508	178.295	132	1.28334	0.609666	32.9641	0.921071	0.712333	41.52	31.4545	33	25	38
120	447.446	371.2	393	1.84933	0.809466	32.9641	0.921071	0.712333	102.17	25.8654	26	20	31
121	435.448	108.209	91	1.42731	0.753533	30.7661	0.911801	0.707123	126.17	41.3626	43	33	51
122	431.589	245.641	131	1.47057	0.811488	32.9149	0.911136	0.782778	42.81	32.7023	34	25	39
123	444.345	472.391	110	1.28412	0.725103	33.0515	0.923237	0.691723	41.98	25.2697	27	20	30
124	447.904	87.9959	178	1.34618	0.755103	31.8945	0.923237	0.691723	41.98	38.8091	40	29	47
125	457.904	87.9959	178	1.34618	0.755103	31.8945	0.923237	0.691723	41.98	38.8091	40	29	47
126	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
127	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
128	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
129	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
130	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
131	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
132	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
133	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
134	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
135	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
136	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
137	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
138	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
139	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
140	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
141	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
142	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
143	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
144	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
145	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
146	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
147	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
148	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
149	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
150	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
151	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
152	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
153	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
154	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
155	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
156	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
157	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
158	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
159	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
160	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
161	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
162	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
163	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
164	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
165	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
166	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
167	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
168	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
169	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
170	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
171	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
172	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
173	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
174	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
175	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
176	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
177	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
178	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
179	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
180	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
181	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
182	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
183	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
184	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
185	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
186	456.785	408.54	113	1.31601	0.688112	31.9948	0.911667	0.807143	43.63	38.4104	39	30	48
187	456.785	408.54	113	1.31601	0.688112	31.9948	0.91						

EV Table 1.doc

1	1	132.831	416.005	118	1.37941	0.71603	12.2573	0.979134	0.7375	4370	37.0339	39	29	44
2	1	141.089	178.484	192	1.45987	0.84491	15.4733	0.91466	0.671431	4755	24.7456	25	19	31
3	1	146.775	477.285	298	1.51175	0.932975	18.4753	0.872504	0.575573	5110	31.9128	33	25	39
4	1	149.47	318.468	161	1.53501	0.959744	13.2598	0.815584	0.721077	4314	30.5957	32	24	37
5	1	151.921	36.2411	203	1.55944	0.976085	18.0148	0.841484	0.763138	5182	45.2315	47	35	55
6	1	153.379	219.534	189	1.58944	0.980532	15.2726	0.815644	0.75	4903	25.9418	26	20	32
7	1	155.872	91.6746	169	1.61944	0.985132	14.4064	0.821192	0.616035	4619	27.5089	28	21	34
8	1	157.379	139.7168	135	1.64944	0.989744	13.4704	0.821192	0.616035	4619	40.4866	42	31	49
9	1	159.872	139.7168	135	1.67944	0.994356	12.5344	0.821192	0.616035	4619	31.7976	33	25	39
10	1	162.379	388.369	198	1.70944	0.998968	11.5984	0.821192	0.616035	4619	26.1394	27	20	32
11	1	164.872	501.03	336	1.73944	0.998968	10.6624	0.821192	0.616035	4619	29.2798	31	22	34
12	1	167.379	160.975	160	1.76944	0.998968	9.7264	0.821192	0.616035	4619	27.975	29	22	34
13	1	169.872	312.893	177	1.79944	0.998968	8.7904	0.821192	0.616035	4619	34.4407	36	31	41
14	1	172.379	312.893	177	1.82944	0.998968	7.8544	0.821192	0.616035	4619	40.2851	44	34	53
15	1	174.872	312.893	177	1.85944	0.998968	6.9184	0.821192	0.616035	4619	30.2851	31	24	34
16	1	177.379	145.944	113	1.88944	0.998968	5.9824	0.821192	0.616035	4619	31.1577	32	26	39
17	1	179.872	436.351	331	1.91944	0.998968	5.0464	0.821192	0.616035	4619	28.1577	29	26	39
18	1	182.379	254.049	308	1.94944	0.998968	4.1104	0.821192	0.616035	4619	28.1577	29	26	39
19	1	184.872	466.48	152	1.97944	0.998968	3.1744	0.821192	0.616035	4619	28.1577	29	26	39
20	1	187.379	354.361	208	2.00944	0.998968	2.2384	0.821192	0.616035	4619	28.1577	29	26	39
21	1	189.872	312.893	177	2.03944	0.998968	1.3024	0.821192	0.616035	4619	28.1577	29	26	39
22	1	192.379	312.893	177	2.06944	0.998968	0.3664	0.821192	0.616035	4619	28.1577	29	26	39
23	1	194.872	312.893	177	2.09944	0.998968	-0.5696	0.821192	0.616035	4619	28.1577	29	26	39
24	1	197.379	312.893	177	2.12944	0.998968	-1.4936	0.821192	0.616035	4619	28.1577	29	26	39
25	1	199.872	312.893	177	2.15944	0.998968	-2.4176	0.821192	0.616035	4619	28.1577	29	26	39
26	1	202.379	312.893	177	2.18944	0.998968	-3.3416	0.821192	0.616035	4619	28.1577	29	26	39
27	1	204.872	312.893	177	2.21944	0.998968	-4.2656	0.821192	0.616035	4619	28.1577	29	26	39
28	1	207.379	312.893	177	2.24944	0.998968	-5.1896	0.821192	0.616035	4619	28.1577	29	26	39
29	1	209.872	312.893	177	2.27944	0.998968	-6.1136	0.821192	0.616035	4619	28.1577	29	26	39
30	1	212.379	312.893	177	2.30944	0.998968	-7.0376	0.821192	0.616035	4619	28.1577	29	26	39
31	1	214.872	312.893	177	2.33944	0.998968	-7.9616	0.821192	0.616035	4619	28.1577	29	26	39
32	1	217.379	312.893	177	2.36944	0.998968	-8.8856	0.821192	0.616035	4619	28.1577	29	26	39
33	1	219.872	312.893	177	2.39944	0.998968	-9.8096	0.821192	0.616035	4619	28.1577	29	26	39
34	1	222.379	312.893	177	2.42944	0.998968	-10.7336	0.821192	0.616035	4619	28.1577	29	26	39
35	1	224.872	312.893	177	2.45944	0.998968	-11.6576	0.821192	0.616035	4619	28.1577	29	26	39
36	1	227.379	312.893	177	2.48944	0.998968	-12.5816	0.821192	0.616035	4619	28.1577	29	26	39
37	1	229.872	312.893	177	2.51944	0.998968	-13.5056	0.821192	0.616035	4619	28.1577	29	26	39
38	1	232.379	312.893	177	2.54944	0.998968	-14.4296	0.821192	0.616035	4619	28.1577	29	26	39
39	1	234.872	312.893	177	2.57944	0.998968	-15.3536	0.821192	0.616035	4619	28.1577	29	26	39
40	1	237.379	312.893	177	2.60944	0.998968	-16.2776	0.821192	0.616035	4619	28.1577	29	26	39
41	1	239.872	312.893	177	2.63944	0.998968	-17.2016	0.821192	0.616035	4619	28.1577	29	26	39
42	1	242.379	312.893	177	2.66944	0.998968	-18.1256	0.821192	0.616035	4619	28.1577	29	26	39
43	1	244.872	312.893	177	2.69944	0.998968	-19.0496	0.821192	0.616035	4619	28.1577	29	26	39
44	1	247.379	312.893	177	2.72944	0.998968	-19.9736	0.821192	0.616035	4619	28.1577	29	26	39
45	1	249.872	312.893	177	2.75944	0.998968	-20.8976	0.821192	0.616035	4619	28.1577	29	26	39
46	1	252.379	312.893	177	2.78944	0.998968	-21.8216	0.821192	0.616035	4619	28.1577	29	26	39
47	1	254.872	312.893	177	2.81944	0.998968	-22.7456	0.821192	0.616035	4619	28.1577	29	26	39
48	1	257.379	312.893	177	2.84944	0.998968	-23.6696	0.821192	0.616035	4619	28.1577	29	26	39
49	1	259.872	312.893	177	2.87944	0.998968	-24.5936	0.821192	0.616035	4619	28.1577	29	26	39
50	1	262.379	312.893	177	2.90944	0.998968	-25.5176	0.821192	0.616035	4619	28.1577	29	26	39
51	1	264.872	312.893	177	2.93944	0.998968	-26.4416	0.821192	0.616035	4619	28.1577	29	26	39
52	1	267.379	312.893	177	2.96944	0.998968	-27.3656	0.821192	0.616035	4619	28.1577	29	26	39
53	1	269.872	312.893	177	2.99944	0.998968	-28.2896	0.821192	0.616035	4619	28.1577	29	26	39
54	1	272.379	312.893	177	3.02944	0.998968	-29.2136	0.821192	0.616035	4619	28.1577	29	26	39
55	1	274.872	312.893	177	3.05944	0.998968	-30.1376	0.821192	0.616035	4619	28.1577	29	26	39
56	1	277.379	312.893	177	3.08944	0.998968	-31.0616	0.821192	0.616035	4619	28.1577	29	26	39
57	1	279.872	312.893	177	3.11944	0.998968	-31.9856	0.821192	0.616035	4619	28.1577	29	26	39
58	1	282.379	312.893	177	3.14944	0.998968	-32.9096	0.821192	0.616035	4619	28.1577	29	26	39
59	1	284.872	312.893	177	3.17944	0.998968	-33.8336	0.821192	0.616035	4619	28.1577	29	26	39
60	1	287.379	312.893	177	3.20944	0.998968	-34.7576	0.821192	0.616035	4619	28.1577	29	26	39
61	1	289.872	312.893	177	3.23944	0.998968	-35.6816	0.821192	0.616035	4619	28.1577	29	26	39
62	1	292.379	312.893	177	3.26944	0.998968	-36.6056	0.821192	0.616035	4619	28.1577	29	26	39
63	1	294.872	312.893	177	3.29944	0.998968	-37.5296	0.821192	0.616035	4619	28.1577	29	26	39
64	1	297.379	312.893	177	3.32944	0.998968	-38.4536	0.821192	0.616035	4619	28.1577	29	26	39
65	1	299.872	312.893	177	3.35944	0.998968	-39.3776	0.821192	0.616035	4619	28.1577	29	26	39
66	1	302.379	312.893	177	3.38944	0.998968	-40.3016	0.821192	0.616035	4619	28.1577	29	26	39
67	1	304.872	312.893	177	3.41944	0.998968	-41.2256	0.821192	0.616035	4619	28.1577	29	26	39
68	1	307.379	312.893	177	3.44944	0.998968	-42.1496	0.821192	0.616035	4619	28.1577	29	26	39
69	1	309.872	312.893	177	3.47944	0.998968	-43.0736	0.821192	0.616035	4619	28.1577	29	26	39
70	1	312.379	312.893	177	3.50944	0.998968	-43.9976	0.821192	0.616035	4619	28.1577	29	26	39
71	1	314.872	312.893	177	3.53944	0.998968	-44.9216	0.821192	0.616035	4619	28.1577	29	26	39
72	1	317.379	312.893	177	3.56944	0.998968	-45.8456	0.821192	0.616035	4619	28.1577	29	26	39
73	1	319.872	312.893	177	3.59944	0.998968	-46.7696	0.821192	0.616035	4619	28.1577	29	26	39
74	1	322.379	312.893	177	3.62944	0.998968	-47.6936	0.821192	0.616035	4619	28.1577	29	26	39
75	1	324.872	312.893	177	3.65944	0.998968	-48.6176	0.821192	0.616035	4619	28.1577	29	26	39
76	1	327.379	312.893	177	3.68944	0.998968	-49.5416	0.821192	0.616035	4619	28.1577	29	26	39
77	1	329.872	312.893	177	3.71944	0.998968	-50.4656	0.821192	0.616035	4619	28.1577	29	26	39
78	1	332.379	312.893	177	3.74944	0.998968	-51.3896	0.821192	0.616035	4619	28.1577	29	26	39
79	1	334.872	312.893	177	3.77944	0.998968	-52.3136	0.821192	0.616035	4619	28.1577	29	26	39
80	1	337.379	312.8											

EV Table 1.doc

1	1	11.1154	387.587	108	1.42566	0.712937	11.5073	0.904348	0.533235	3955	34.0286	39	31	46
2	2	9.80103	425.712	66	1.20459	0.559891	9.167	0.88	0.733232	3059	46.3155	48	53	44
3	3	11.4146	443.818	77	1.37652	0.524846	9.90149	0.927711	0.7	2950	38.2312	40	34	44
4	4	14.4143	456.346	70	1.03328	0.23174	9.407	0.860676	0.7	3970	56.7113	56	46	58
5	5	18.3947	455.1316	114	1.263	0.610824	12.0478	0.912	0.710769	3992	34.1404	35	27	41
6	6	21.18	472.59	100	1.52012	0.753156	11.2838	0.884956	0.714286	3840	38.4	39	47	47
7	7	25.1143	499.143	169	1.52652	0.755554	13.3512	0.903216	0.673077	4072	23.0837	30	22	34
8	8	32.3148	432.905	169	1.95543	0.859145	15.5126	0.936283	0.63	12608	64.709	69	81	84
9	9	37.4847	450.614	114	1.27031	0.616685	12.0478	0.919355	0.707169	1004	35.1278	37	28	43
10	10	47.468	418.544	125	1.47033	0.721089	12.6157	0.905797	0.694644	7138	57.104	60	44	49
11	11	57.3023	425.684	132	1.23156	0.56462	13.0131	0.917243	0.738849	4708	31.6391	32	25	39
12	12	68.4594	444.607	112	1.60845	0.86045	11.9848	0.91129	0.724759	4075	36.0619	37	29	44
13	13	80.7057	480.193	196	1.01853	0.31803	15.7973	0.903226	0.608811	5557	28.352	30	22	35
14	14	94.7162	311.425	196	1.07457	0.374947	13.4935	0.923581	0.785714	4063	28.4126	29	23	34
15	15	109.0119	311.425	252	1.28906	0.61066	17.9125	0.927427	0.572727	11427	45.3452	45	38	58
16	16	125.7822	423.473	132	1.10508	0.45066	12.4334	0.917293	0.782051	1642	38.2131	39	30	46
17	17	146.7934	400.731	134	1.60428	0.78187	13.0419	0.937063	0.681673	4152	30.9951	32	25	37
18	18	167.566	404.566	212	1.49816	0.716224	16.4294	0.917749	0.69281	4804	41.3283	42	32	52
19	19	201.3013	464.719	92	1.47318	0.722855	10.822	0.910881	0.69697	3641	39.5761	39	31	49
20	20	240.733	464.719	478	1.03578	0.240563	24.67	0.910476	0.735385	14880	31.7297	32	24	38
21	21	284.6333	328.808	120	1.21491	0.567192	12.3408	0.895532	0.710058	4070	31.9167	35	27	41
22	22	341.703	370.181	221	1.5985	0.780155	16.7746	0.934641	0.734667	7891	35.7059	37	28	43
23	23	414.7356	329.558	156	1.54808	0.743771	14.0935	0.927077	0.718789	4298	27.5513	28	21	34
24	24	464.2813	40.0764	144	2.23108	0.93326	13.5406	0.917368	0.8	5206	38.1528	37	28	44
25	25	505.8089	208.4	91	1.33133	0.621074	10.8817	0.902513	0.65035	3739	40.2043	41	33	49
26	26	571.2843	193.915	117	1.42001	0.704191	12.2053	0.905917	0.709091	3961	33.8547	31	26	41
27	27	624.8938	104.974	149	2.01167	0.87453	15.5126	0.917416	0.678287	5541	28.3175	30	23	36
28	28	74.4616	178.771	134	1.70184	0.89451	13.0619	0.911565	0.697917	4152	30.8851	33	24	37
29	29	84.4636	109.166	107	1.23867	0.590113	11.672	0.90678	0.716252	4231	38.5234	39	30	43
30	30	85.2385	148.102	117	2.7253	0.90841	13.6809	0.90184	0.660192	4168	28.7823	29	23	36
31	31	84.232	167.611	113	1.16561	0.537718	11.9948	0.90184	0.724359	4168	38.8052	38	29	46
32	32	90.8089	208.4	102	1.6489	0.743068	22.6321	0.846619	0.649553	14872	38.9032	37	28	45
33	33	90.5396	160.101	109	1.37451	0.727416	11.7808	0.892443	0.61881	4053	37.1835	39	30	45
34	34	98.5058	463.579	273	1.51649	0.852109	18.6328	0.877816	0.59684	13955	51.1172	52	38	64
35	35	99.5127	247.923	259	1.62483	0.788136	19.5115	0.937304	0.684211	9270	32.0067	33	25	39
36	36	98.2744	177.536	277	2.96801	0.82225	18.7817	0.882168	0.507232	7730	27.3061	29	21	34
37	37	98.7634	166.432	93	1.67058	0.801152	16.3559	0.885714	0.65035	3656	41.4674	42	32	50
38	38	108.312	493.449	205	2.11642	0.787276	16.3559	0.911111	0.711808	4628	22.5756	23	17	28
39	39	132.24	441.38	150	2.11642	0.787276	16.3559	0.911111	0.711808	4628	22.5756	23	17	28
40	40	131.82	222.161	117	1.92467	0.787276	16.3559	0.911111	0.711808	4628	22.5756	23	17	28
41	41	131.82	222.161	117	1.92467	0.787276	16.3559	0.911111	0.711808	4628	22.5756	23	17	28
42	42	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
43	43	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
44	44	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
45	45	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
46	46	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
47	47	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
48	48	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
49	49	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
50	50	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
51	51	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
52	52	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
53	53	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
54	54	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
55	55	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
56	56	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
57	57	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
58	58	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
59	59	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
60	60	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
61	61	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
62	62	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
63	63	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
64	64	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
65	65	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
66	66	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
67	67	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
68	68	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
69	69	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
70	70	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
71	71	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
72	72	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
73	73	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
74	74	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
75	75	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61
76	76	132.83	411.358	108	1.32867	0.654417	13.2053	0.916377	0.757576	7793	51.9533	53	40	61

EV Table 1.doc	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105	106	107	108	109	110	111	112	113	114	115	116	117	118	119	120	121	122	123	124	125	126	127	128	129	130	131	132	133	134	135	136	137	138	139	140	141	142	143	144	145	146	147	148	149	150	151	152	153	154	155	156	157	158	159	160	161	162	163	164	165	166	167	168	169	170	171	172	173	174	175	176	177	178	179	180	181	182	183	184	185	186	187	188	189	190	191	192	193	194	195	196	197	198	199	200	201	202	203	204	205	206	207	208	209	210	211	212	213	214	215	216	217	218	219	220	221	222	223	224	225	226	227	228	229	230	231	232	233	234	235	236	237	238	239	240	241	242	243	244	245	246	247	248	249	250	251	252	253	254	255	256	257	258	259	260	261	262	263	264	265	266	267	268	269	270	271	272	273	274	275	276	277	278	279	280	281	282	283	284	285	286	287	288	289	290	291	292	293	294	295	296	297	298	299	300	301	302	303	304	305	306	307	308	309	310	311	312	313	314	315	316	317	318	319	320	321	322	323	324	325	326	327	328	329	330	331	332	333	334	335	336	337	338	339	340	341	342	343	344	345	346	347	348	349	350	351	352	353	354	355	356	357	358	359	360	361	362	363	364	365	366	367	368	369	370	371	372	373	374	375	376	377	378	379	380	381	382	383	384	385	386	387	388	389	390	391	392	393	394	395	396	397	398	399	400	401	402	403	404	405	406	407	408	409	410	411	412	413	414	415	416	417	418	419	420	421	422	423	424	425	426	427	428	429	430	431	432	433	434	435	436	437	438	439	440	441	442	443	444	445	446	447	448	449	450	451	452	453	454	455	456	457	458	459	460	461	462	463	464	465	466	467	468	469	470	471	472	473	474	475	476	477	478	479	480	481	482	483	484	485	486	487	488	489	490	491	492	493	494	495	496	497	498	499	500	501	502	503	504	505	506	507	508	509	510	511	512	513	514	515	516	517	518	519	520	521	522	523	524	525	526	527	528	529	530	531	532	533	534	535	536	537	538	539	540	541	542	543	544	545	546	547	548	549	550	551	552	553	554	555	556	557	558	559	560	561	562	563	564	565	566	567	568	569	570	571	572	573	574	575	576	577	578	579	580	581	582	583	584	585	586	587	588	589	590	591	592	593	594	595	596	597	598	599	600	601	602	603	604	605	606	607	608	609	610	611	612	613	614	615	616	617	618	619	620	621	622	623	624	625	626	627	628	629	630	631	632	633	634	635	636	637	638	639	640	641	642	643	644	645	646	647	648	649	650	651	652	653	654	655	656	657	658	659	660	661	662	663	664	665	666	667	668	669	670	671	672	673	674	675	676	677	678	679	680	681	682	683	684	685	686	687	688	689	690	691	692	693	694	695	696	697	698	699	700	701	702	703	704	705	706	707	708	709	710	711	712	713	714	715	716	717	718	719	720	721	722	723	724	725	726	727	728	729	730	731	732	733	734	735	736	737	738	739	740	741	742	743	744	745	746	747	748	749	750	751	752	753	754	755	756	757	758	759	760	761	762	763	764	765	766	767	768	769	770	771	772	773	774	775	776	777	778	779	780	781	782	783	784	785	786	787	788	789	790	791	792	793	794	795	796	797	798	799	800	801	802	803	804	805	806	807	808	809	810	811	812	813	814	815	816	817	818	819	820	821	822	823	824	825	826	827	828	829	830	831	832	833	834	835	836	837	838	839	840	841	842	843	844	845	846	847	848	849	850	851	852	853	854	855	856	857	858	859	860	861	862	863	864	865	866	867	868	869	870	871	872	873	874	875	876	877	878	879	880	881	882	883	884	885	886	887	888	889	890	891	892	893	894	895	896	897	898	899	900	901	902	903	904	905	906	907	908	909	910	911	912	913	914	915	916	917	918	919	920	921	922	923	924	925	926	927	928	929	930	931	932	933	934	935	936	937	938	939	940	941	942	943	944	945	946	947	948	949	950	951	952	953	954	955	956	957	958	959	960	961	962	963	964	965	966	967	968	969	970	971	972	973	974	975	976	977	978	979	980	981	982	983	984	985	986	987	988	989	990	991	992	993	994	995	996	997	998	999	1000
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1	21.2188	364.135	54	1.29161	0.631372	9.07003	0.476712	0.727273	3672	56.9063	37	44	59
2	24.0233	453.946	129	1.42766	0.877231	12.8159	0.921479	0.685536	3672	30.0155	30	23	37
3	35.4428	114.293	140	1.45761	0.727231	12.8159	0.909091	0.727273	4313	30.8071	31	23	37
4	39.9272	251.376	206	1.4719	0.79493	16.1953	0.919613	0.777632	9011	43.7573	35	35	53
5	38.5582	328.843	178	1.06005	0.331735	15.0545	0.927083	0.791111	9196	51.6629	52	38	64
6	38.3667	41.52	150	1.17638	0.577089	13.8198	0.920245	0.765306	4709	31.3933	33	23	37
7	38.3767	362.388	85	1.17668	0.577265	10.4031	0.894737	0.708333	3980	46.4255	49	39	57
8	45.227	24.1727	139	1.40448	0.702172	13.7036	0.894737	0.708333	4278	30.7682	32	23	37
9	45.227	101.968	187	1.77163	0.825128	15.4304	0.912193	0.697857	4410	23.5849	24	18	25
10	53.8904	62.5278	146	1.61579	0.785475	13.6343	0.921031	0.693238	4298	28.4384	30	22	36
11	53.8904	326.379	124	1.31697	0.650318	12.466	0.9	0.693238	4098	32.5238	33	23	36
12	65.4198	170.966	324	2.47732	0.514923	20.3108	0.79803	0.482183	16200	50	52	59	61
13	61.8615	292.138	195	1.45539	0.798019	15.1571	0.919811	0.711286	9025	66.7821	58	55	61
14	61.8615	214.022	180	1.7726	0.825678	13.1388	0.904323	0.711286	8730	48.5	50	37	59
15	73.2222	335.178	45	1.47858	0.711415	7.5694	0.919367	0.711286	3366	78.9	74	56	74
16	77.3137	227.052	153	1.46508	0.770839	13.9573	0.921487	0.73	4772	29.2289	20	54	59
17	76.2015	362.453	53	1.10916	0.492654	6.21672	0.913793	0.731611	3514	66.2019	69	50	60
18	81.9478	94.3146	192	1.27679	0.621756	15.6353	0.923077	0.761905	8805	45.8594	47	37	43
19	86.0609	317.826	115	1.25501	0.601218	12.1005	0.912698	0.761905	4048	35.2	36	27	43
20	98.0964	190.528	197	1.34408	0.496308	15.4376	0.920561	0.724265	9240	66.4036	48	36	59
21	98.4796	427.551	101	1.32402	0.638326	11.4518	0.895632	0.72028	4108	39.8835	61	31	49
22	102.493	121.67	203	2.8412	0.318412	16.0769	0.815391	0.461564	8738	43.0443	61	31	53
23	100.531	36.639	122	1.72624	0.815018	12.4636	0.913793	0.621449	3920	32.1511	33	28	39
24	101.027	97.8931	145	1.86167	0.810234	13.7736	0.925466	0.622232	4349	29.1819	30	22	33
25	93.8	243.069	170	1.27409	0.611208	12.8455	0.902778	0.714286	4111	31.9308	35	27	41
26	104.422	795.355	109	1.17019	0.519167	11.7806	0.915964	0.756944	4199	38.5729	39	30	46
27	104.422	74.3317	90	1.51552	0.73064	15.9177	0.929907	0.765395	8907	44.7588	47	34	54
28	104.422	441.211	108	1.20505	0.62565	10.7047	0.913157	0.693208	3953	43.9222	45	33	53
29	104.422	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
30	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
31	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
32	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
33	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
34	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
35	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
36	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
37	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
38	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
39	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
40	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
41	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
42	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
43	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
44	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
45	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
46	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
47	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
48	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
49	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
50	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
51	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
52	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
53	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
54	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
55	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
56	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
57	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
58	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
59	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
60	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
61	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
62	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
63	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
64	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
65	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
66	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
67	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
68	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
69	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
70	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
71	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
72	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
73	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
74	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
75	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
76	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
77	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
78	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
79	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
80	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
81	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44
82	116.122	349.443	115	1.22263	0.575349	12.1005	0.912698	0.761905	4134	35.9478	37	28	44

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1	1	9	283.446	101.911	127	1.6816	0.405787	12.7162	0.913669	0.661458	4675	36.4173	27	29	44
2	1	9	284.645	100.611	173	1.51854	0.755961	14.1415	0.925134	0.759772	7670	46.3153	37	35	54
3	1	9	285.844	99.311	219	1.35534	0.758111	10.2318	0.902226	0.661454	4311	50.131	31	39	62
4	1	9	286.043	98.011	265	1.19214	0.759716	14.0028	0.916647	0.733333	4326	29.3696	30	33	36
5	1	9	287.242	96.711	311	1.02894	0.761321	10.7047	0.909091	0.743002	4134	45.9331	46	36	56
6	1	9	288.441	95.411	357	0.86574	0.762926	10.7047	0.896534	0.746086	4067	38.0093	39	29	46
7	1	9	289.640	94.111	403	0.70254	0.764531	11.674	0.884018	0.749170	4000	44.3396	44	36	54
8	1	9	290.839	92.811	449	0.53934	0.766136	13.7273	0.871503	0.752153	3933	60.4527	60	41	74
9	1	9	292.038	91.511	495	0.37614	0.767741	16.0182	0.859019	0.755136	3866	85.4545	88	65	106
10	1	9	293.237	90.211	541	0.21294	0.769346	18.3137	0.846514	0.758119	3799	111.2768	113	92	150
11	1	9	294.436	88.911	587	0.04974	0.770951	20.6092	0.834014	0.761102	3732	137.099	139	121	194
12	1	9	295.635	87.611	633	0.11654	0.772556	22.9047	0.821514	0.764085	3665	162.921	165	143	238
13	1	9	296.834	86.311	679	0.28334	0.774161	25.1902	0.809014	0.767068	3598	188.743	191	165	282
14	1	9	298.033	85.011	725	0.45014	0.775766	27.4757	0.796514	0.770051	3531	214.565	217	187	326
15	1	9	299.232	83.711	771	0.61694	0.777371	29.7612	0.784014	0.773034	3464	240.387	243	209	370
16	1	9	300.431	82.411	817	0.78374	0.778976	32.0467	0.771514	0.776017	3397	266.209	269	231	414
17	1	9	301.630	81.111	863	0.95054	0.780581	34.3322	0.759014	0.779000	3330	292.031	295	253	458
18	1	9	302.829	79.811	909	1.11734	0.782186	36.6177	0.746514	0.781983	3263	317.853	321	275	502
19	1	9	304.028	78.511	955	1.28414	0.783791	38.9032	0.734014	0.784966	3196	343.675	347	297	546
20	1	9	305.227	77.211	1001	1.45094	0.785396	41.1887	0.721514	0.787949	3129	369.497	373	319	590
21	1	9	306.426	75.911	1047	1.61774	0.787001	43.4742	0.709014	0.790932	3062	395.319	399	341	634
22	1	9	307.625	74.611	1093	1.78454	0.788606	45.7597	0.696514	0.793915	2995	421.141	425	363	678
23	1	9	308.824	73.311	1139	1.95134	0.790211	48.0452	0.684014	0.796898	2928	446.963	451	385	722
24	1	9	310.023	72.011	1185	2.11814	0.791816	50.3307	0.671514	0.799881	2861	472.785	477	407	766
25	1	9	311.222	70.711	1231	2.28494	0.793421	52.6162	0.659014	0.802864	2794	498.607	503	429	810
26	1	9	312.421	69.411	1277	2.45174	0.795026	54.9017	0.646514	0.805847	2727	524.429	529	451	854
27	1	9	313.620	68.111	1323	2.61854	0.796631	57.1872	0.634014	0.808830	2660	550.251	555	473	898
28	1	9	314.819	66.811	1369	2.78534	0.798236	59.4727	0.621514	0.811813	2593	576.073	581	495	942
29	1	9	316.018	65.511	1415	2.95214	0.799841	61.7582	0.609014	0.814796	2526	601.895	607	517	986
30	1	9	317.217	64.211	1461	3.11894	0.801446	64.0437	0.596514	0.817779	2459	627.717	633	539	1030
31	1	9	318.416	62.911	1507	3.28574	0.803051	66.3292	0.584014	0.820762	2392	653.539	661	561	1074
32	1	9	319.615	61.611	1553	3.45254	0.804656	68.6147	0.571514	0.823745	2325	679.361	687	583	1118
33	1	9	320.814	60.311	1599	3.61934	0.806261	70.9002	0.559014	0.826728	2258	705.183	715	605	1162
34	1	9	322.013	59.011	1645	3.78614	0.807866	73.1857	0.546514	0.829711	2191	731.005	739	627	1206
35	1	9	323.212	57.711	1691	3.95294	0.809471	75.4712	0.534014	0.832694	2124	756.827	765	649	1250
36	1	9	324.411	56.411	1737	4.11974	0.811076	77.7567	0.521514	0.835677	2057	782.649	791	671	1294
37	1	9	325.610	55.111	1783	4.28654	0.812681	80.0422	0.509014	0.838660	1990	808.471	819	693	1338
38	1	9	326.809	53.811	1829	4.45334	0.814286	82.3277	0.496514	0.841643	1923	834.293	843	715	1382
39	1	9	328.008	52.511	1875	4.62014	0.815891	84.6132	0.484014	0.844626	1856	860.115	871	737	1426
40	1	9	329.207	51.211	1921	4.78694	0.817496	86.8987	0.471514	0.847609	1789	885.937	899	759	1470
41	1	9	330.406	49.911	1967	4.95374	0.819101	89.1842	0.459014	0.850592	1722	911.759	923	781	1514
42	1	9	331.605	48.611	2013	5.12054	0.820706	91.4697	0.446514	0.853575	1655	937.581	949	803	1558
43	1	9	332.804	47.311	2059	5.28734	0.822311	93.7552	0.434014	0.856558	1588	963.403	975	825	1602
44	1	9	334.003	46.011	2105	5.45414	0.823916	96.0407	0.421514	0.859541	1521	989.225	1001	847	1646
45	1	9	335.202	44.711	2151	5.62094	0.825521	98.3262	0.409014	0.862524	1454	1015.047	1027	869	1690
46	1	9	336.401	43.411	2197	5.78774	0.827126	100.6117	0.396514	0.865507	1387	1040.869	1053	891	1734
47	1	9	337.600	42.111	2243	5.95454	0.828731	102.8972	0.384014	0.868490	1320	1066.691	1079	913	1778
48	1	9	338.799	40.811	2289	6.12134	0.830336	105.1827	0.371514	0.871473	1253	1092.513	1105	935	1822
49	1	9	340.000	39.511	2335	6.28814	0.831941	107.4682	0.359014	0.874456	1186	1118.335	1131	957	1866
50	1	9	341.199	38.211	2381	6.45494	0.833546	109.7537	0.346514	0.877439	1119	1144.157	1157	979	1910
51	1	9	342.398	36.911	2427	6.62174	0.835151	112.0392	0.334014	0.880422	1052	1169.979	1183	1001	1954
52	1	9	343.597	35.611	2473	6.78854	0.836756	114.3247	0.321514	0.883405	985	1195.801	1209	1023	1998
53	1	9	344.796	34.311	2519	6.95534	0.838361	116.6102	0.309014	0.886388	918	1221.623	1235	1045	2042
54	1	9	345.995	33.011	2565	7.12214	0.839966	118.8957	0.296514	0.889371	851	1247.445	1261	1067	2086
55	1	9	347.194	31.711	2611	7.28894	0.841571	121.1812	0.284014	0.892354	784	1273.267	1287	1089	2130
56	1	9	348.393	30.411	2657	7.45574	0.843176	123.4667	0.271514	0.895337	717	1299.089	1313	1111	2174
57	1	9	349.592	29.111	2703	7.62254	0.844781	125.7522	0.259014	0.898320	650	1324.911	1339	1133	2218
58	1	9	350.791	27.811	2749	7.78934	0.846386	128.0377	0.246514	0.901303	583	1350.733	1365	1155	2262
59	1	9	351.990	26.511	2795	7.95614	0.847991	130.3232	0.234014	0.904286	516	1376.555	1391	1177	2306
60	1	9	353.189	25.211	2841	8.12294	0.849596	132.6087	0.221514	0.907269	449	1402.377	1417	1199	2350
61	1	9	354.388	23.911	2887	8.28974	0.851201	134.8942	0.209014	0.910252	382	1428.199	1443	1221	2394
62	1	9	355.587	22.611	2933	8.45654	0.852806	137.1797	0.196514	0.913235	315	1454.021	1469	1243	2438
63	1	9	356.786	21.311	2979	8.62334	0.854411	139.4652	0.184014	0.916218	248	1479.843	1495	1265	2482
64	1	9	357.985	20.011	3025	8.79014	0.856016	141.7507	0.171514	0.919201	181	1505.665	1521	1287	2526
65	1	9	359.184	18.711	3071	8.95694	0.857621	144.0362	0.159014	0.922184	114	1531.487	1547	1309	2570
66	1	9	360.383	17.411	3117	9.12374	0.859226	146.3217	0.146514	0.925167	47	1557.309	1573	1331	2614
67	1	9	361.582	16.111	3163	9.29054	0.860831	148.6072	0.134014	0.928150	0	1583.131	1599	1353	2658
68	1	9	362.781	14.811	3209	9.45734	0.862436	150.8927	0.121514	0.931133	0	1608.953	1625	1375	2702
69	1	9	363.980	13.511	3255	9.62414	0.864041	153.1782	0.109014	0.934116	0	1634.775	1651	1397	2746
70	1	9	365.179	12.211	3301	9.79094	0.865646	155.4637	0.096514	0.937099	0	1660.597	1677	1419	2790
71	1	9	366.378	10.911	3347	9.95774	0.867251	157.7492	0.084014	0.940082	0	1686.419	1703	1441	2834
72	1	9	367.577	9.611	3393	10.12454	0.868856	160.0347	0.071514	0.943065	0	1712.241	1729	1463	2878
73	1	9	368.776	8.311	3439	10.29134	0.870461	162.3202	0.059014	0.946048	0	1738.063	1755	1485	2922
74	1	9	369.975	7.011	3485	10.45814	0.872066	164.6057	0.046514	0.949031	0	1763.885	1781	1507	2966
75	1	9													

CLAIMS

What is claimed is:

1. A method of predicting a property of a manipulation of cells based
5 upon a descriptor, said method comprising:
 providing a plurality of cells;
 manipulating said plurality of cells;
 capturing a morphological value from said plurality of cells;
 assigning a degree of presence of said morphological value; and
10 storing said morphological value and said degree of presence;
 wherein said descriptor is derived from a first component of a cell and
 a second component of said cell, said capturing said morphometric value from said
 plurality of cells comprises determining a relationship between said first component
 and said second component.
- 15 2. The method of claim 1 wherein said first component and said second
 component are selected from a protein, a protein modification, a nucleic acid, a lipid,
 a carbohydrate, a subcellular structure and an organelle.
3. The method of 1 wherein said step of manipulation occurs in a manner
 selected from a electrical source, a chemical source, a thermal source, a gravitational
20 source, a nuclear source, a temporal source, and a biological source
4. The method of claim 3 wherein said chemical source is selected from a
 pharmacological candidate and a drug screening library.
5. The method of claim 1 wherein said morphological value is selected
 from a count, an area, a perimeter, a length, a breadth, a fiber length, a fiber breadth, a
25 shape factor, a elliptical form factor, an inner radius, an outer radius, a mean radius,
 an equivalent radius, an equivalent sphere volume, an equivalent prolate volume, an
 equivalent oblate volume, an equivalent sphere surface area, an average gray value, a
 total gray value, and an optical density.
6. The method of claim 1 wherein said degree of presence is
30 multiple of a quantized value.

7. A computer program product for populating a database with manipulated biological information, said computer program product comprising:
- code for providing a plurality of cells in various stages of the cell cycle, said stages of the cell cycle including at least one selected from interphase, G0 phase, G1 phase, S phase, G2 phase, M phase, prophase, prometaphase, metaphase, anaphase, and telophase;
 - code for manipulating said cells in said various stages of cell cycle development to form a plurality of manipulated cells;
 - code for capturing an image of said plurality of manipulated cells;
 - code for determining a descriptor from said image for said manipulated cells;
 - code for populating a database with said descriptor;
 - wherein said image includes a first component of a cell and a second component of said cell; and
 - a computer readable storage medium for holding the codes.
8. The computer program product of claim 7 wherein said first component and said second component are selected from a protein, a protein modification, a nucleic acid, a lipid, a carbohydrate, a sub-cellular structure and an organelle.
9. The computer program product of claim 7 wherein said image is a digitized representation of said plurality of manipulated cells.
11. The computer program product of claim 9 wherein said digitized representation provides a density value of said plurality of manipulated cells.
11. The computer program product of claim 7 wherein said descriptors comprise numeric or logical values.
12. The computer program product of claim 11 wherein said values further comprises a nucleotide.
13. The computer program product of claim 11 wherein said values further comprises an amino acid letter.
14. A system for capturing images of cells or cell structures, the system comprising:
- a cell holder comprising a plurality of sites in a spatial orientation, each of the sites being capable of holding a plurality of cells to be imaged;

an image capturing device coupled to the cell holder, the image capture device being adapted to capture at least one image in at least one of the plurality of sites;

an illumination apparatus comprising a liquid light guide coupled to the plate for highlighting the plurality of cells in a relatively even spatial manner for image capturing purposes;

an image processing device coupled to the image capturing device, the image capturing device being adapted to convert the image into a digital representation; and

a database storage device comprising a database management element coupled to the image capturing device, the database storage device being adapted to retrieve the digital representation of the image from the image processing device and storing the digital representation.

15. The system of claim 14 further comprising a stage comprising a device for moving the cell holder in a spatial direction to traverse across the cell holder in the spatial orientation.

16. The system of claim 14 wherein the illumination apparatus comprises sub-elements, at least one of the sub-elements being positioned away from the image capturing device to prevent a possibility of vibration from the one sub-elements to be transmitted to the image capturing device.

17. The system of claim 14 wherein the digital representation comprises a plurality of regions and objects.

18. The system of claim 14 further comprising a computing device connected between the database storage device and the image processing device.

19. The system of claim 14 wherein the image capturing device comprises a magnification of at least 1X and greater to capture the image of the site.

20. The system of claim 14 wherein the plurality of sites comprises at least 96 sites.

21. The system of claim 14 wherein the liquid light guide characterized as a flexible member that substantially prevents vibration from the an element of the illumination apparatus to be transferred to the image capturing device.

22. The system of claim 14 wherein the spatial direction can be selected from an x-direction, a y-direction, or a z-direction in a Cartesian coordinate system.

23. The system of claim 14 wherein the each of the sites comprises
5 a volume that is sufficient to prevent a solution therein from evaporating in a substantial manner that may influence the image capturing.

24. A method for identifying a mechanism of action for a first compound, the method comprising the steps of:
receiving the first compound;
10 measuring at least one feature of a cellular phenotype to define a target phenotype;
identifying additional compounds providing a feature similar to the feature identified in the measuring step; and
characterizing the first compound in terms of distance from a specific
15 target phenotype having known characteristics.

25. The method of claim 24 comprising the further step of storing the additional compounds and their associated phenotypes in a database.

26. A method for identifying a mechanism of action for a cellular stimulus, the method comprising the steps of:
20 receiving cells of interest;
measuring at least one feature of the cells to define and quantify a target phenotype;
identifying additional compounds providing a feature similar to the feature identified in the measuring step; and
25 characterizing the first compound in terms of distance from a specific target phenotype having known characteristics.

27. A method for identifying information relevant to at least one of a mechanism of action and cellular activity by utilizing assay data to elucidate a phenotype, the method comprising the steps of:
30 identifying a target protein;
identifying positive and negative biochemical hits related to the target protein;
defining the target phenotype utilizing the positive and negative hits;

identifying other compounds providing similar features; and
characterizing the first compound in terms of distance from a specific
target phenotype having known characteristics.

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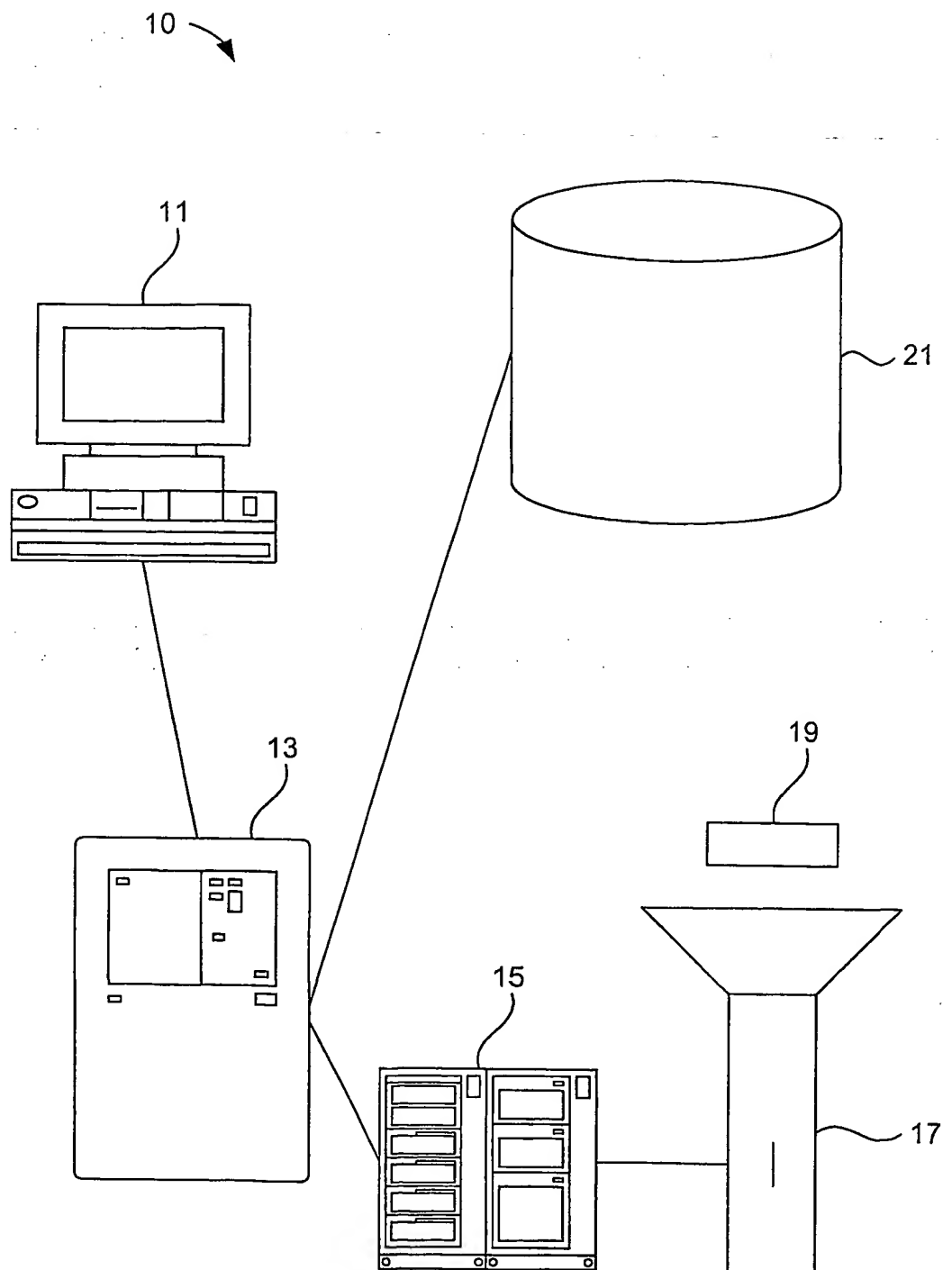


FIG. 1

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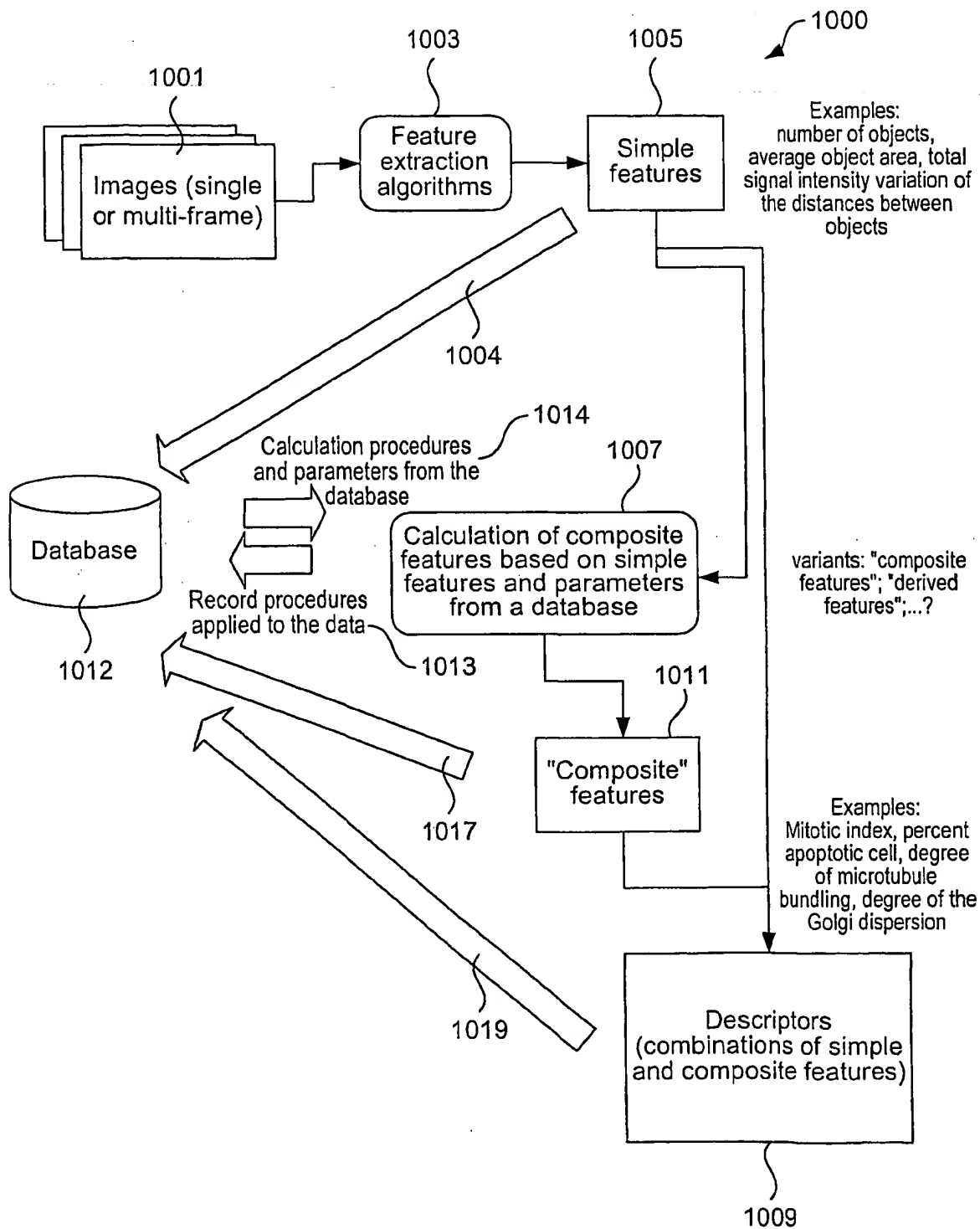


FIG. 1A

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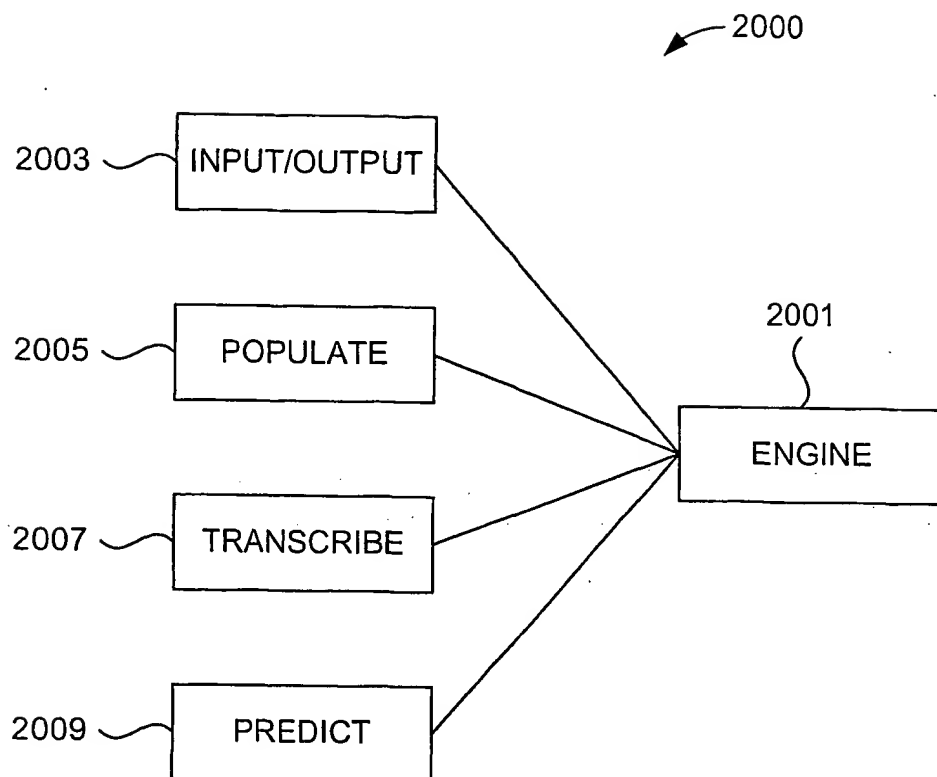


FIG. 1B

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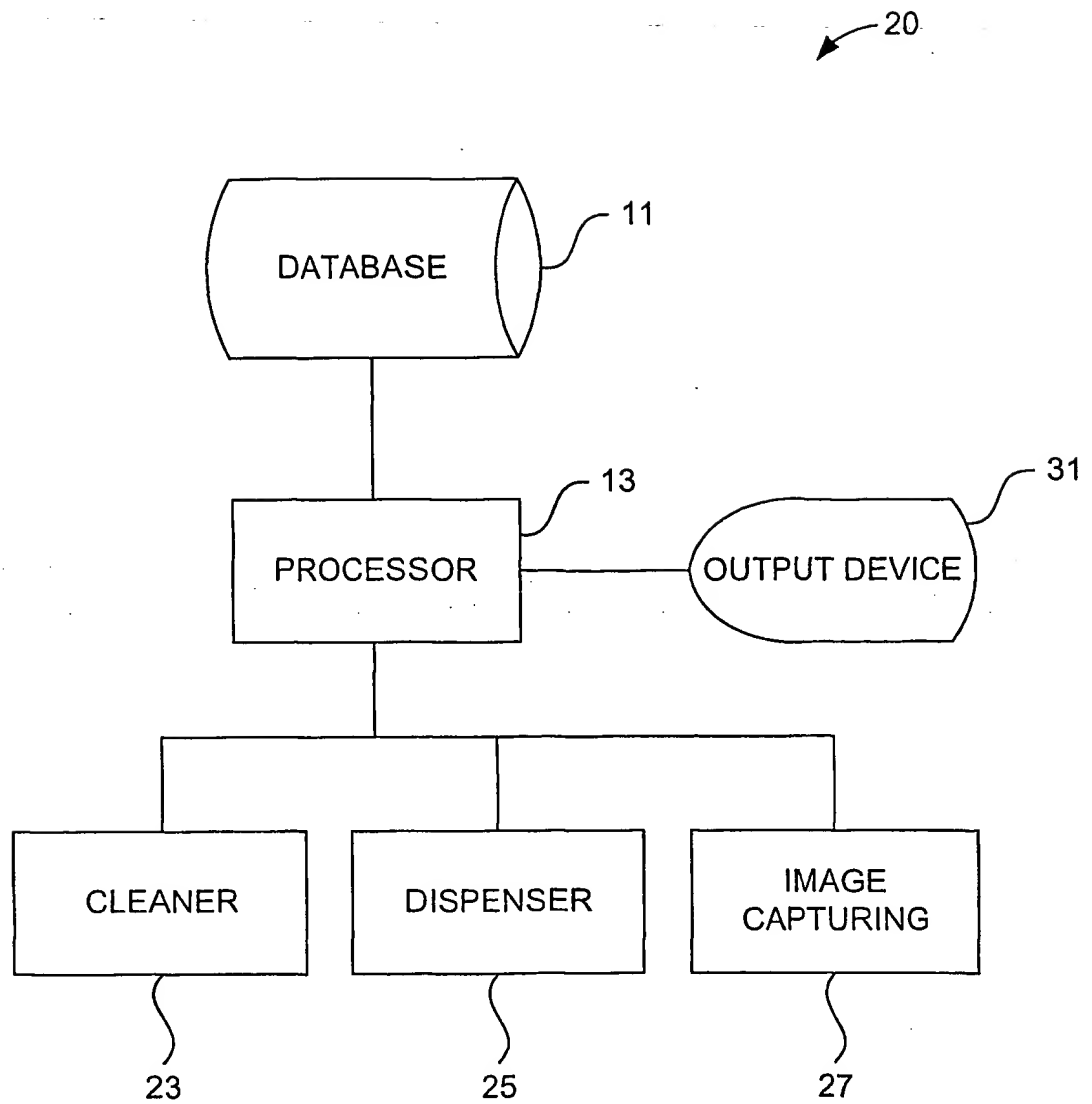


FIG. 2

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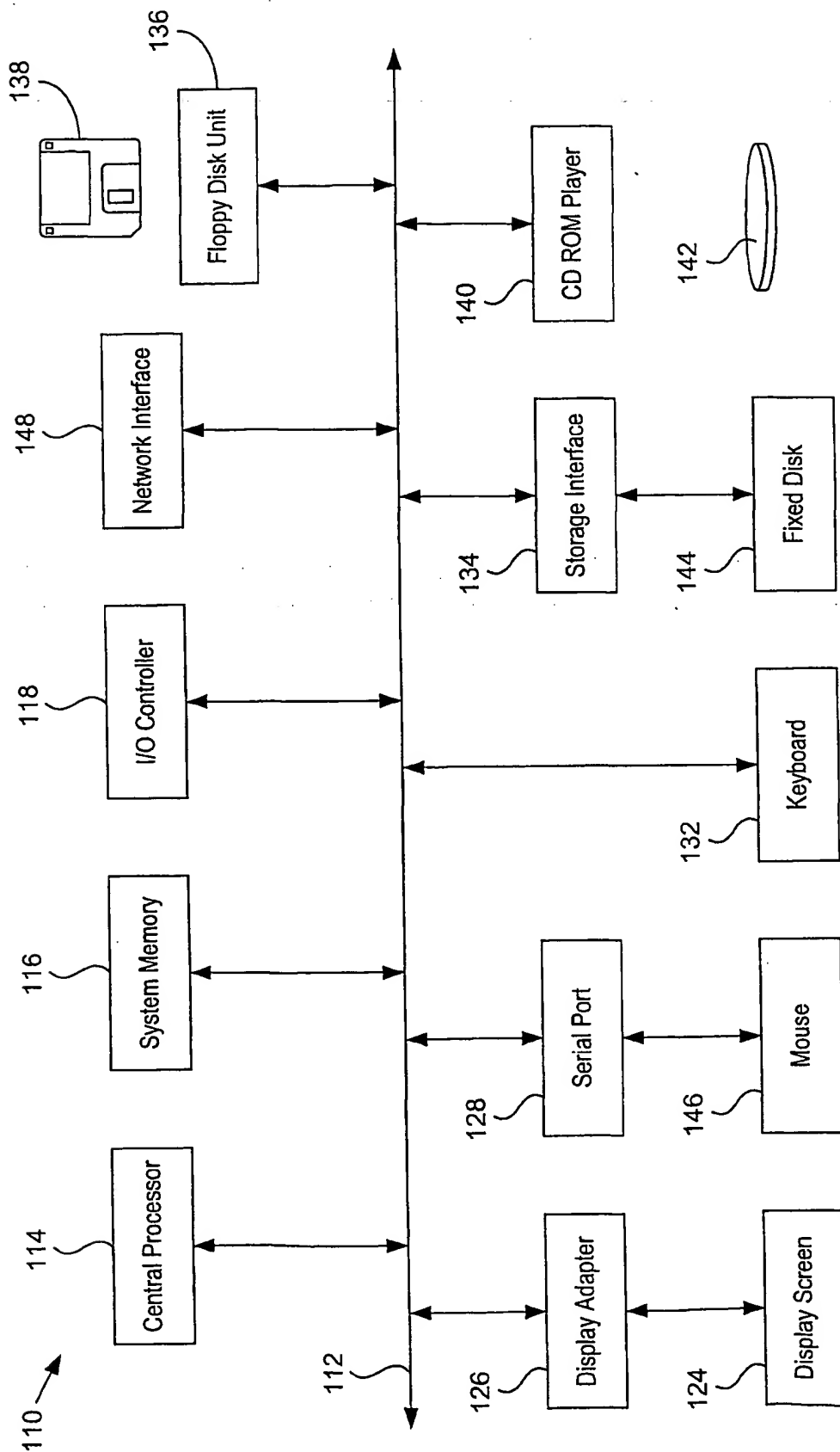


FIG. 3

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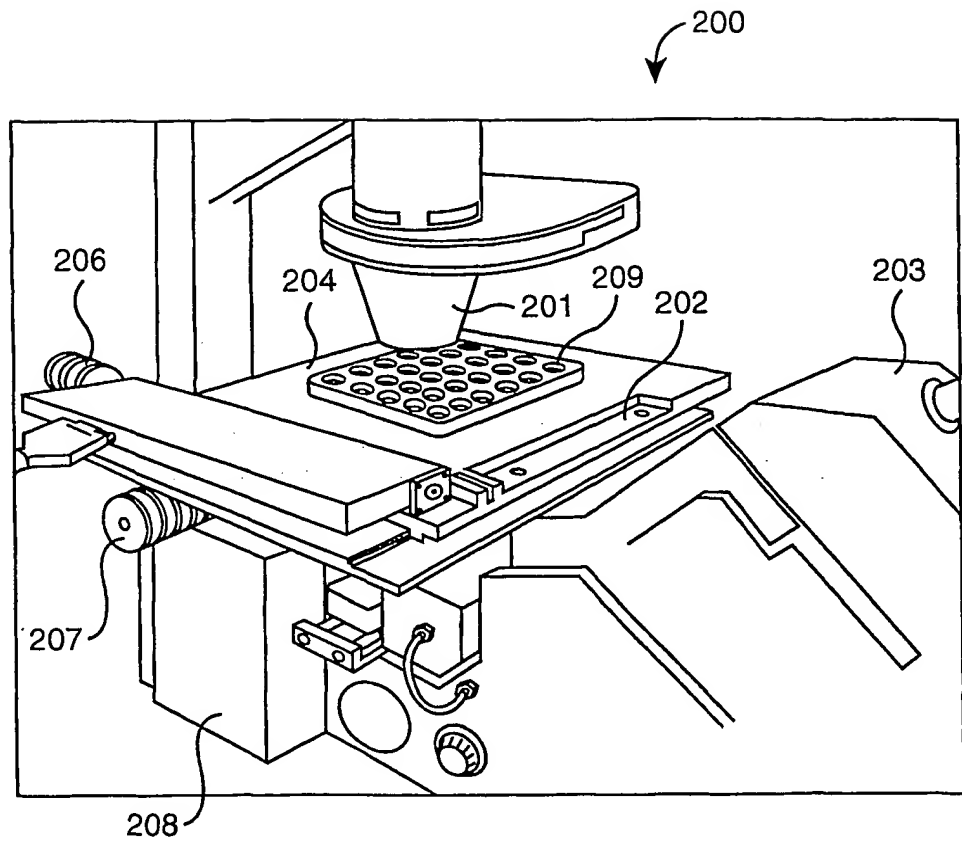


FIG. 4

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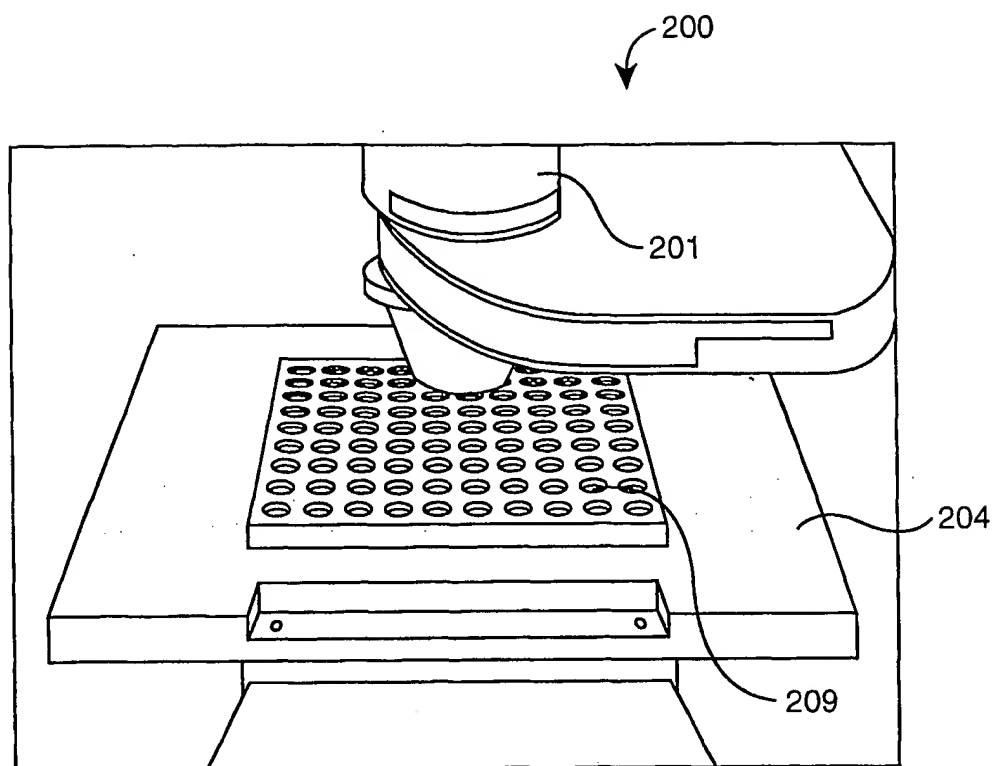


FIG. 5

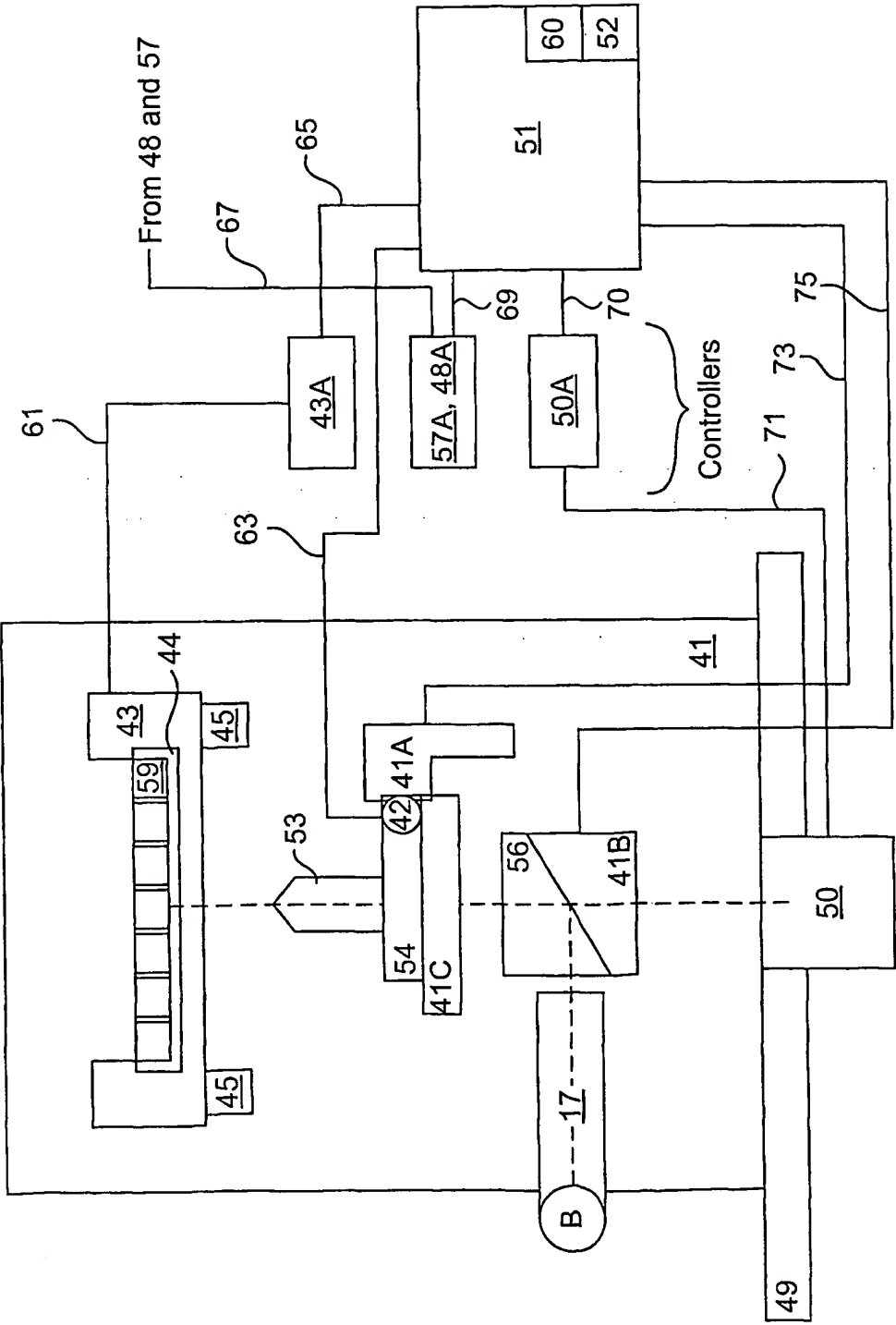


FIG. 5A

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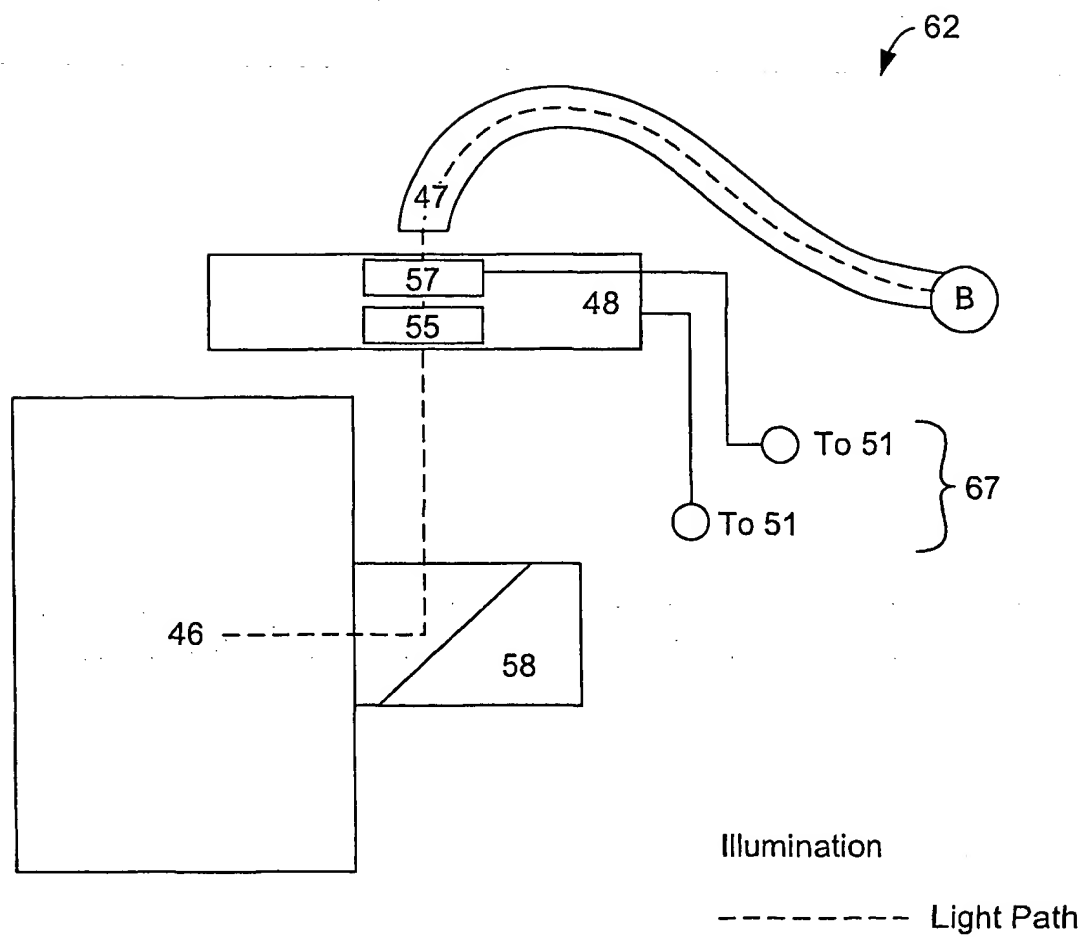


FIG. 5B

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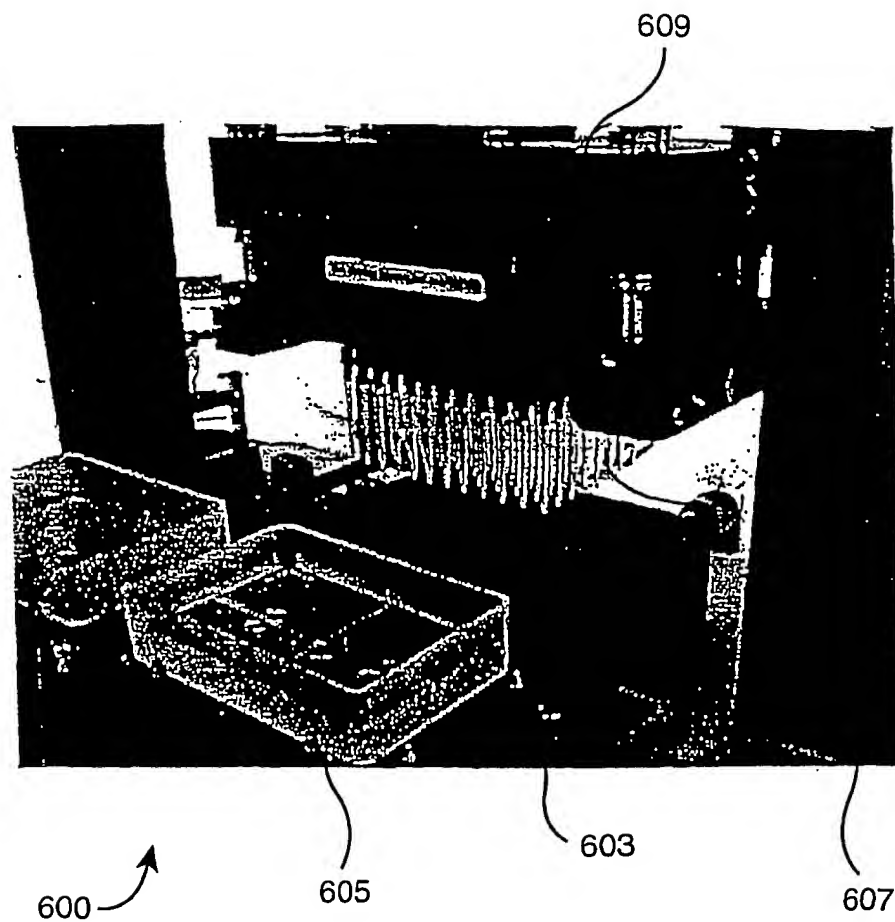


FIG. 6

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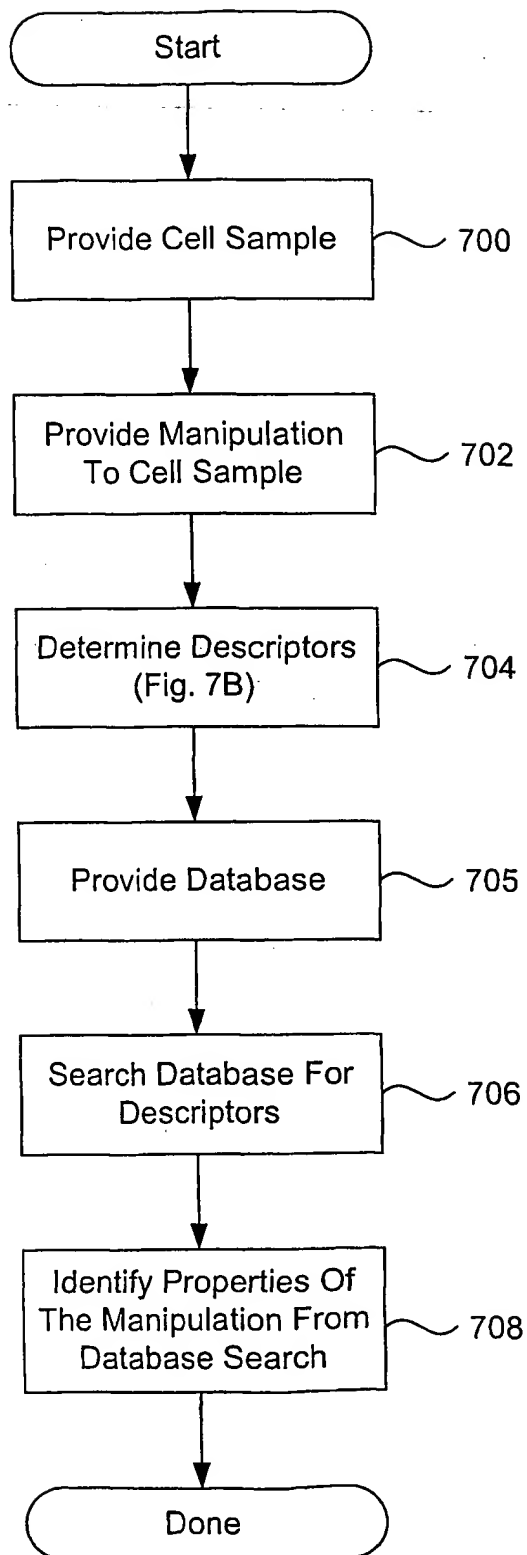


FIG. 7A

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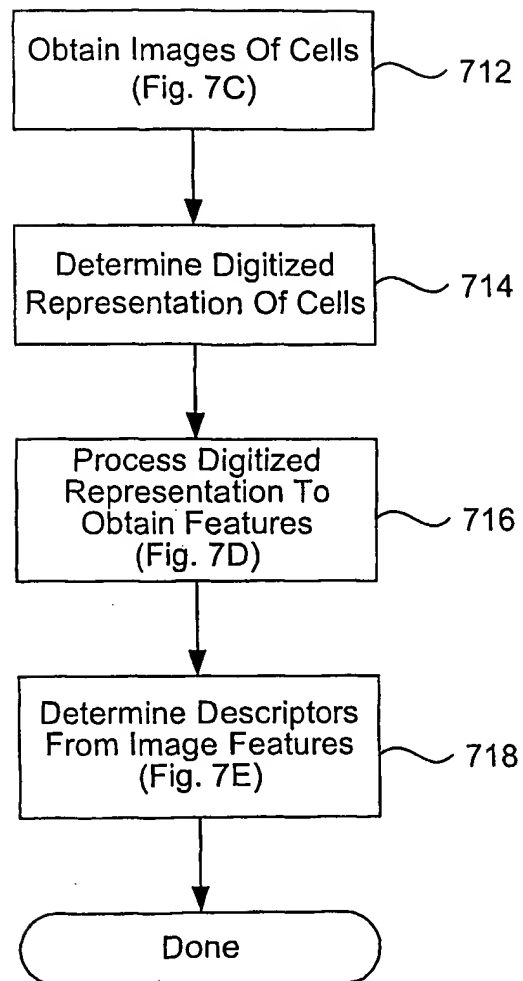


FIG. 7B
Step 704 of Fig. 7A

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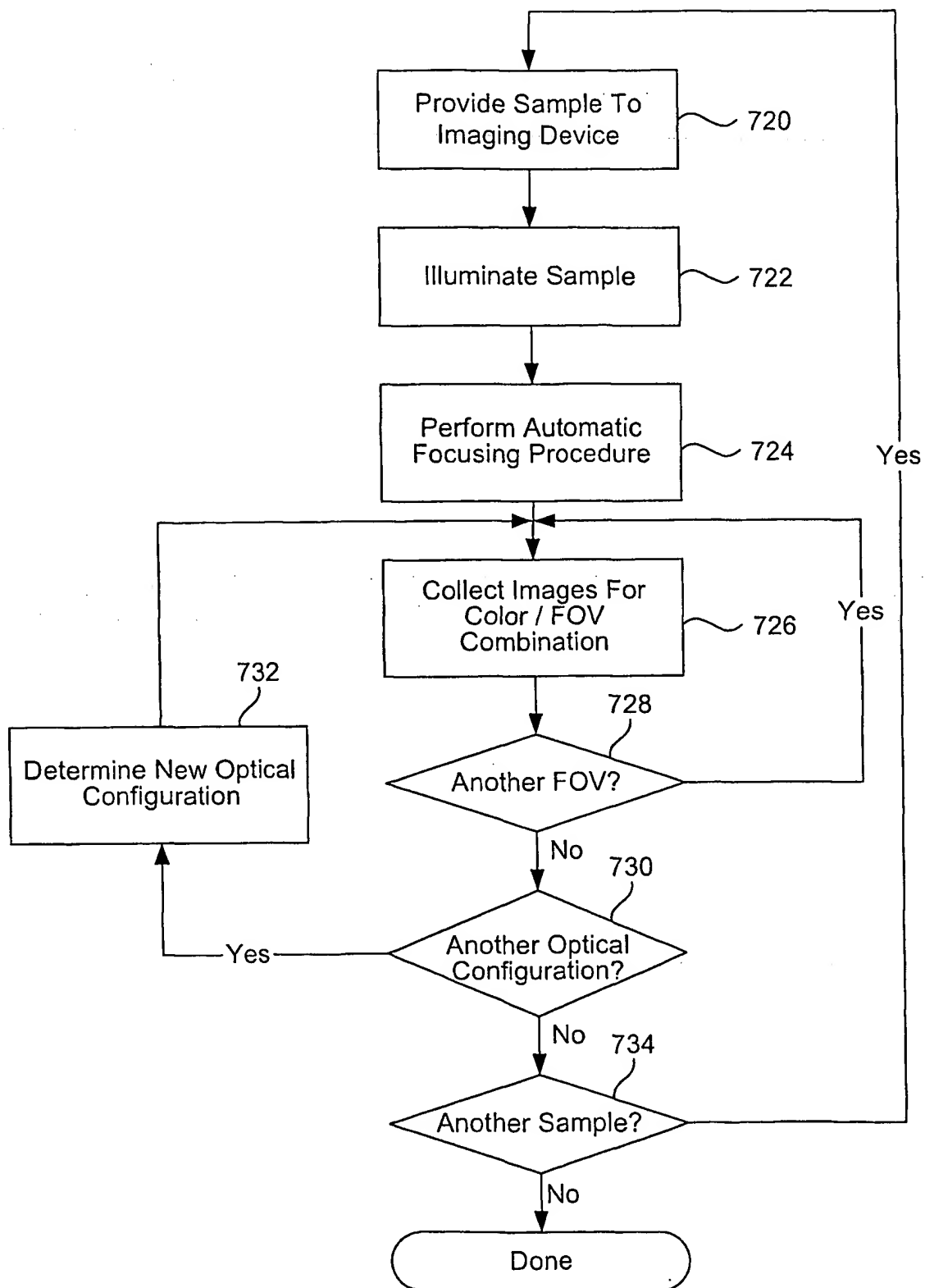


FIG. 7C

Step 714 of Fig. 7B

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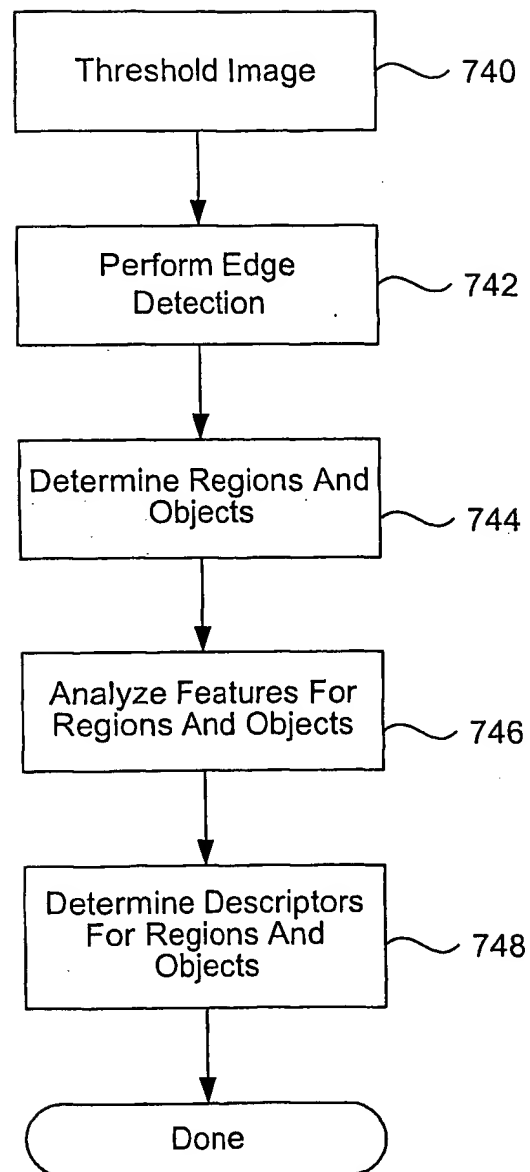


FIG. 7D
Step 716 of Fig. 7B

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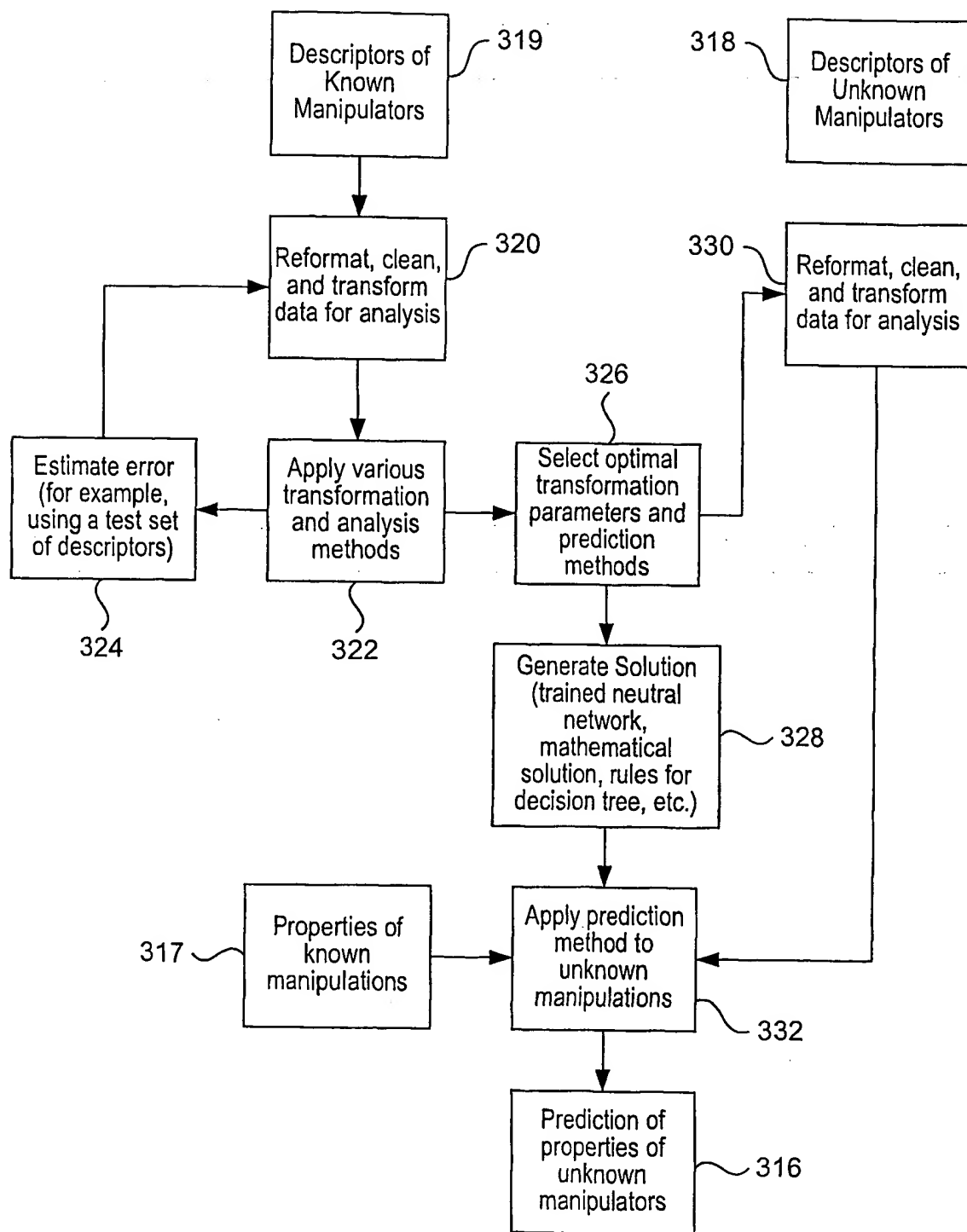


FIG. 7E

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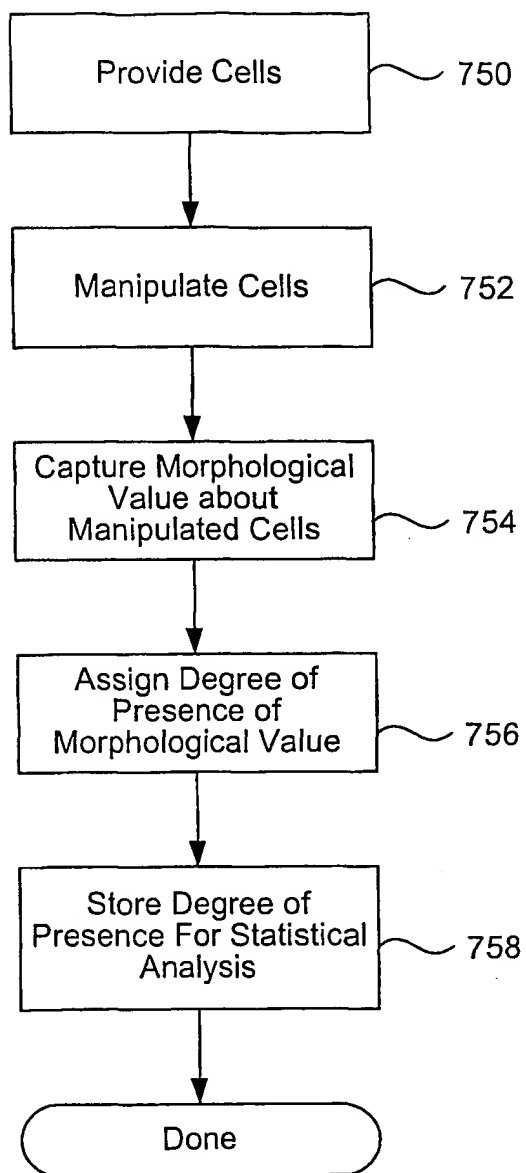


FIG. 7F

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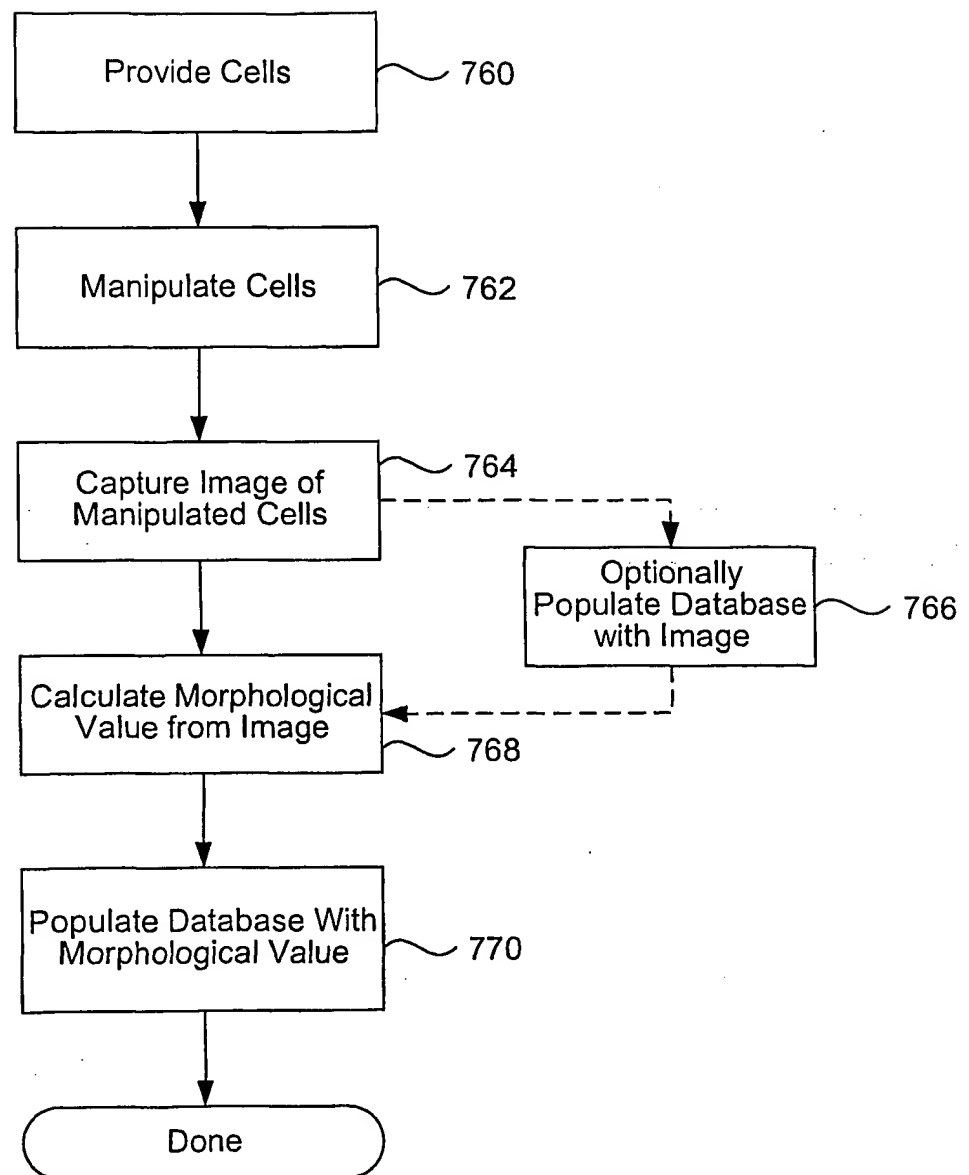


FIG. 7G

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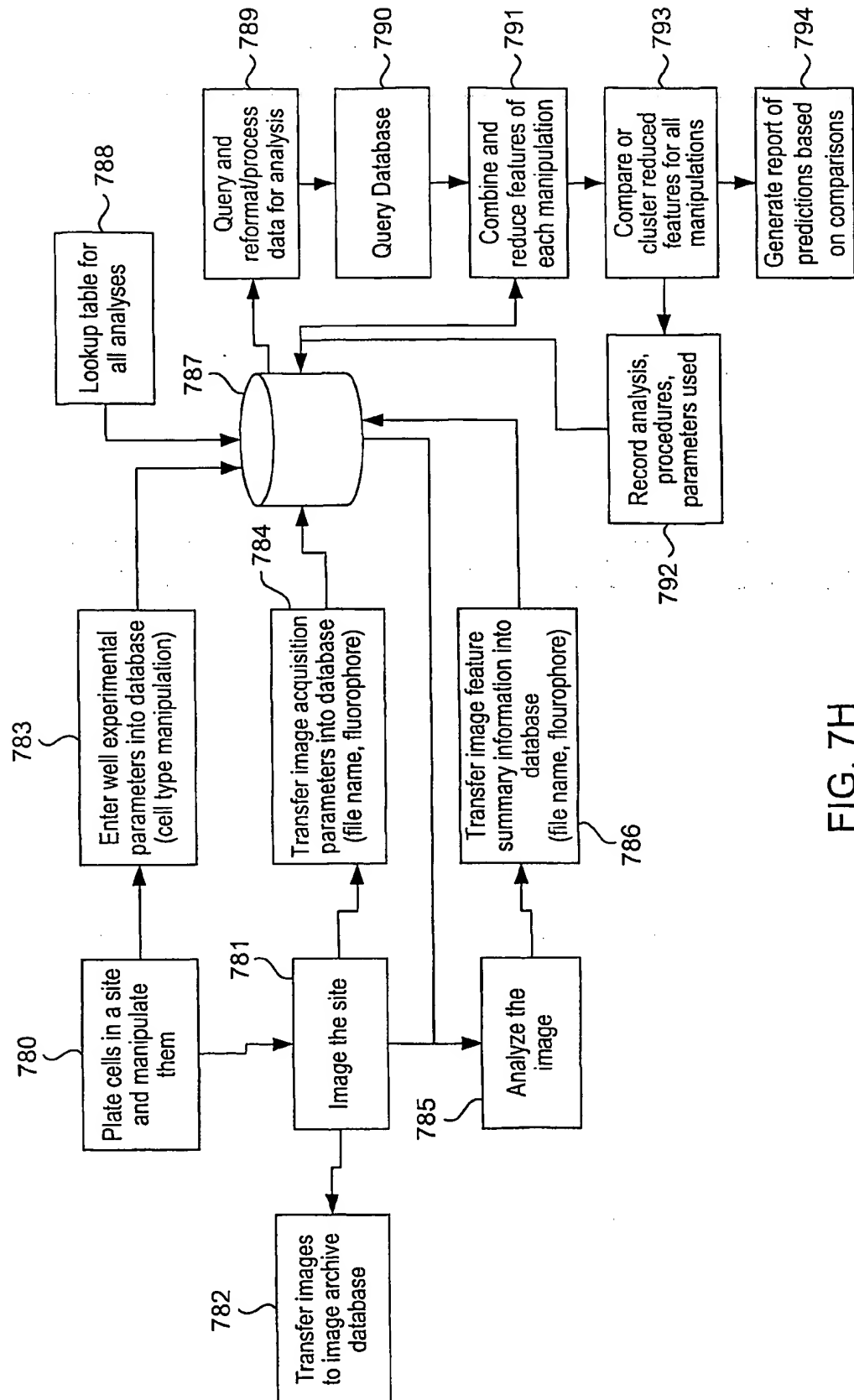


FIG. 7H

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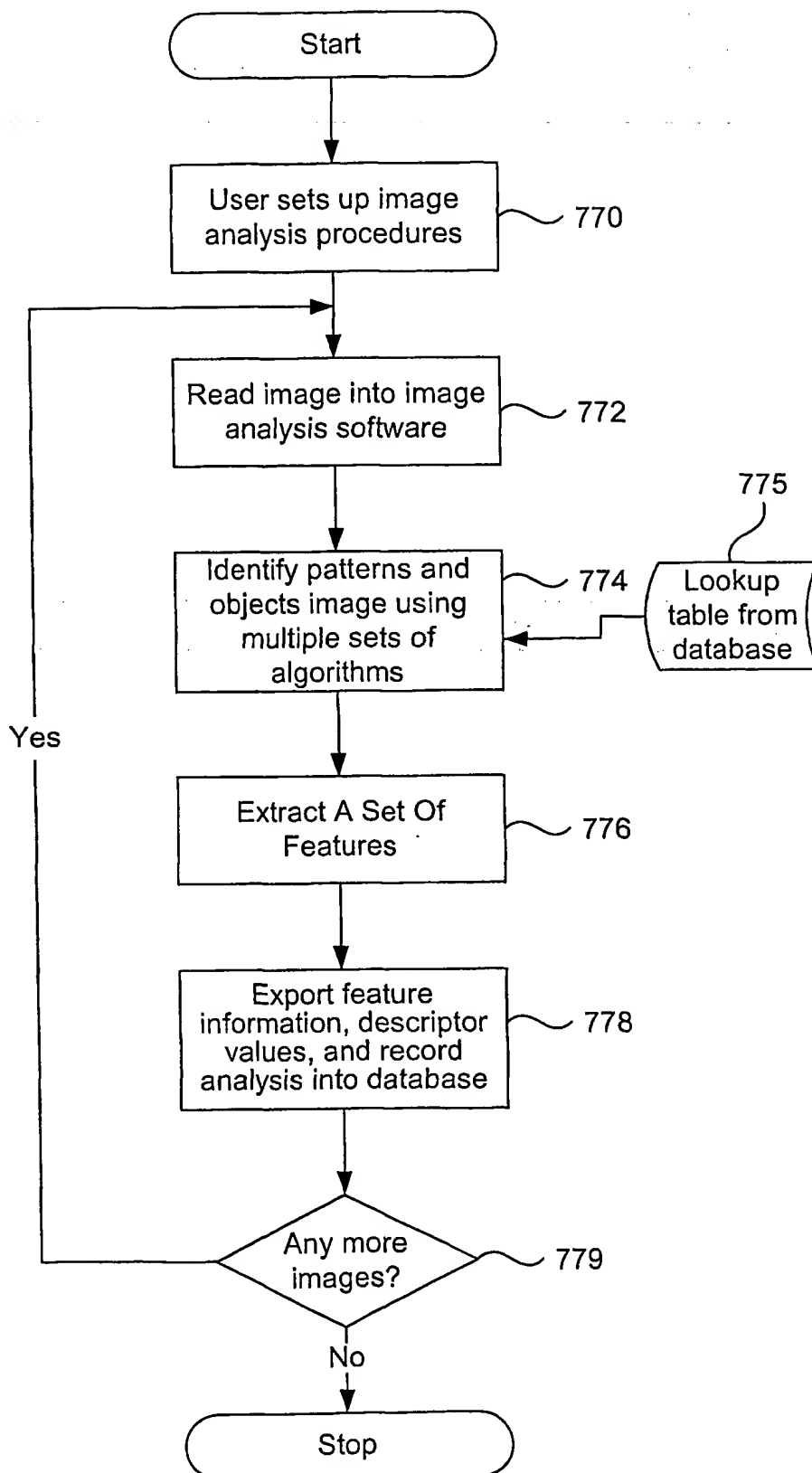


FIG. 71

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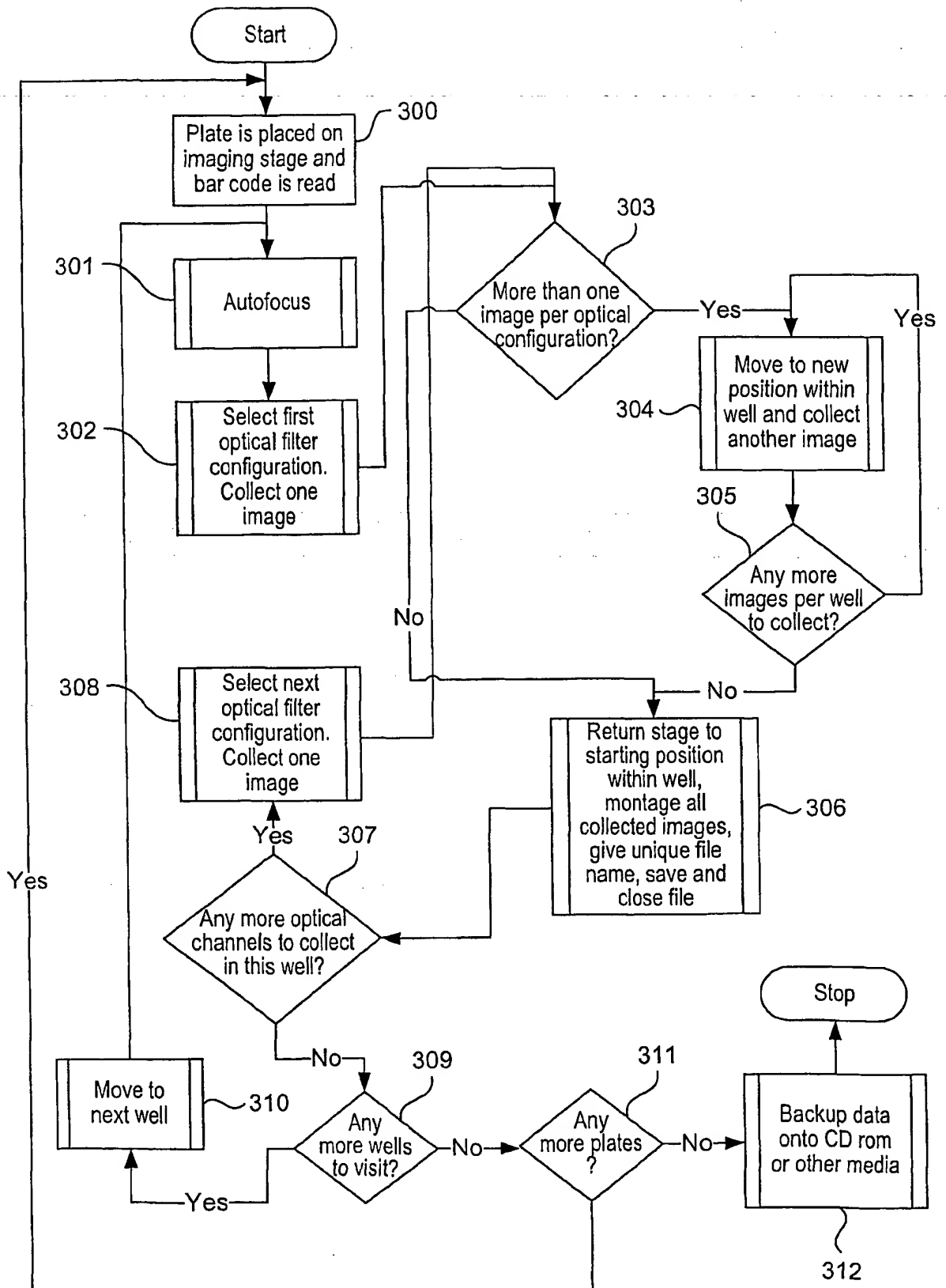


FIG. 7J

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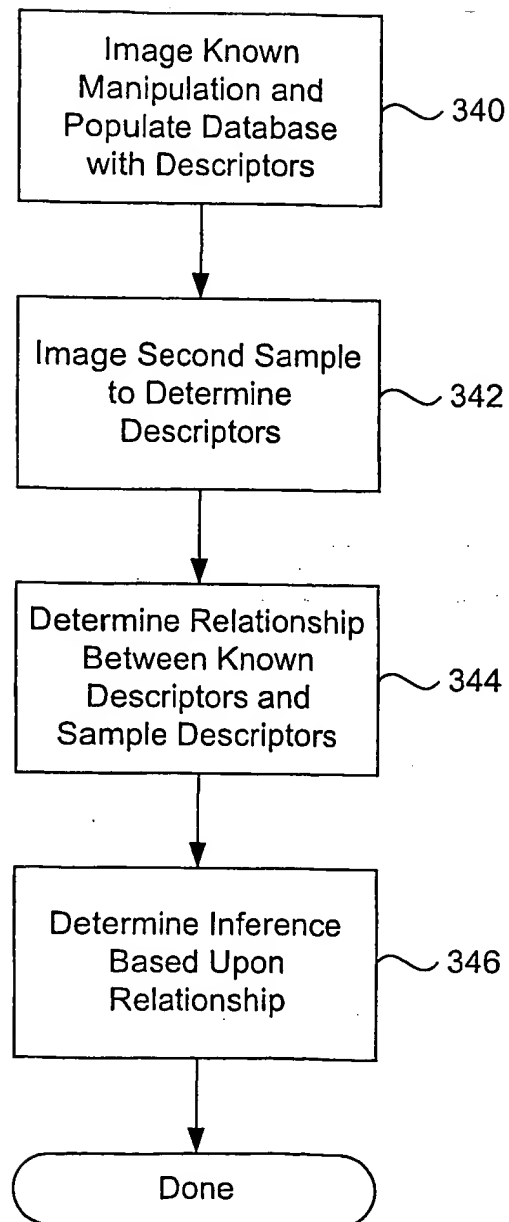


FIG. 7K

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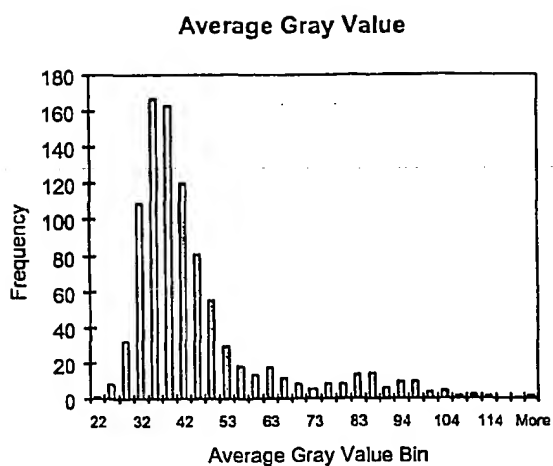


FIG. 8A

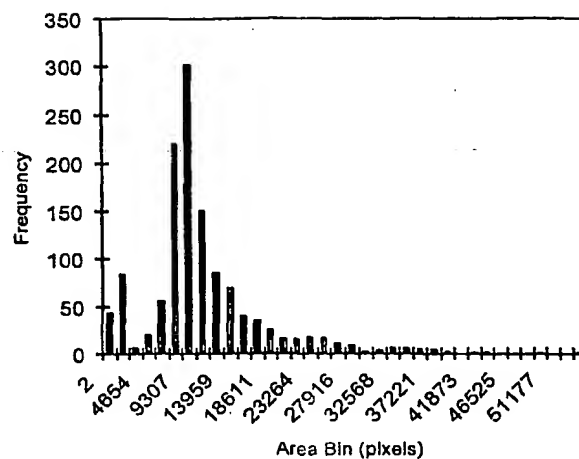


FIG. 8B

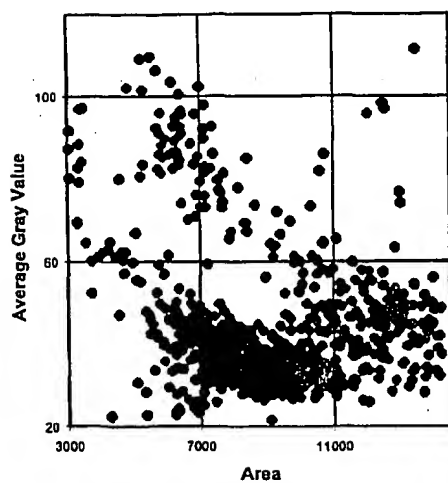


FIG. 8C

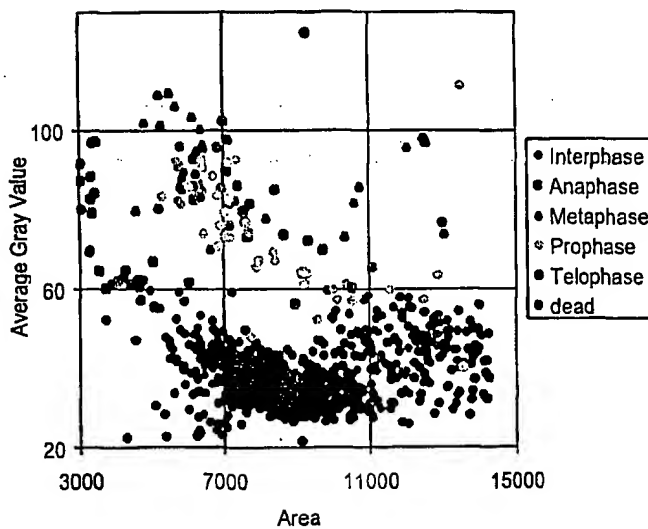


FIG. 8D

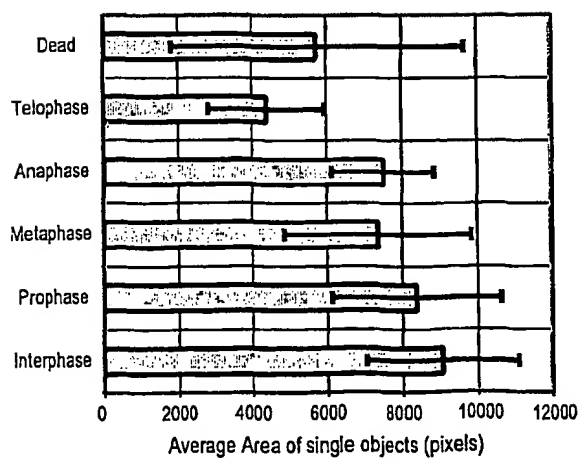


FIG. 8E

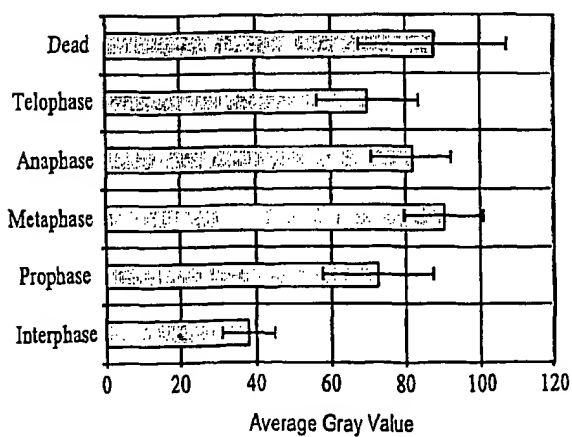


FIG. 8F

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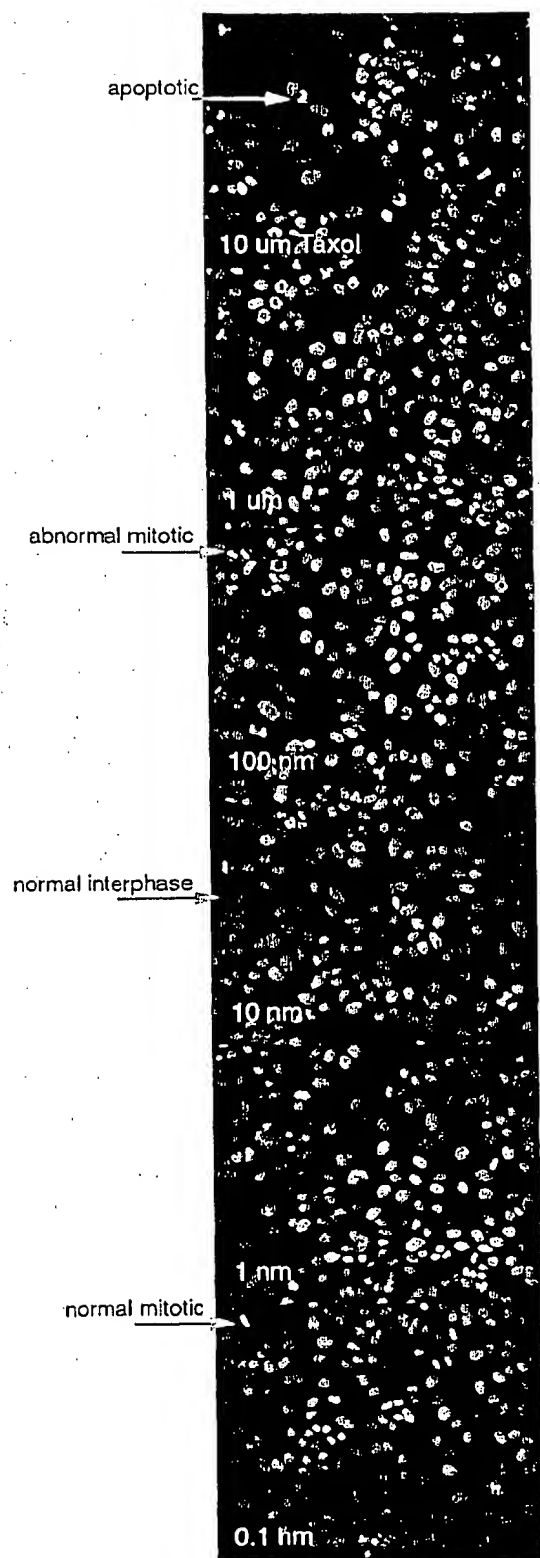


FIG. 9

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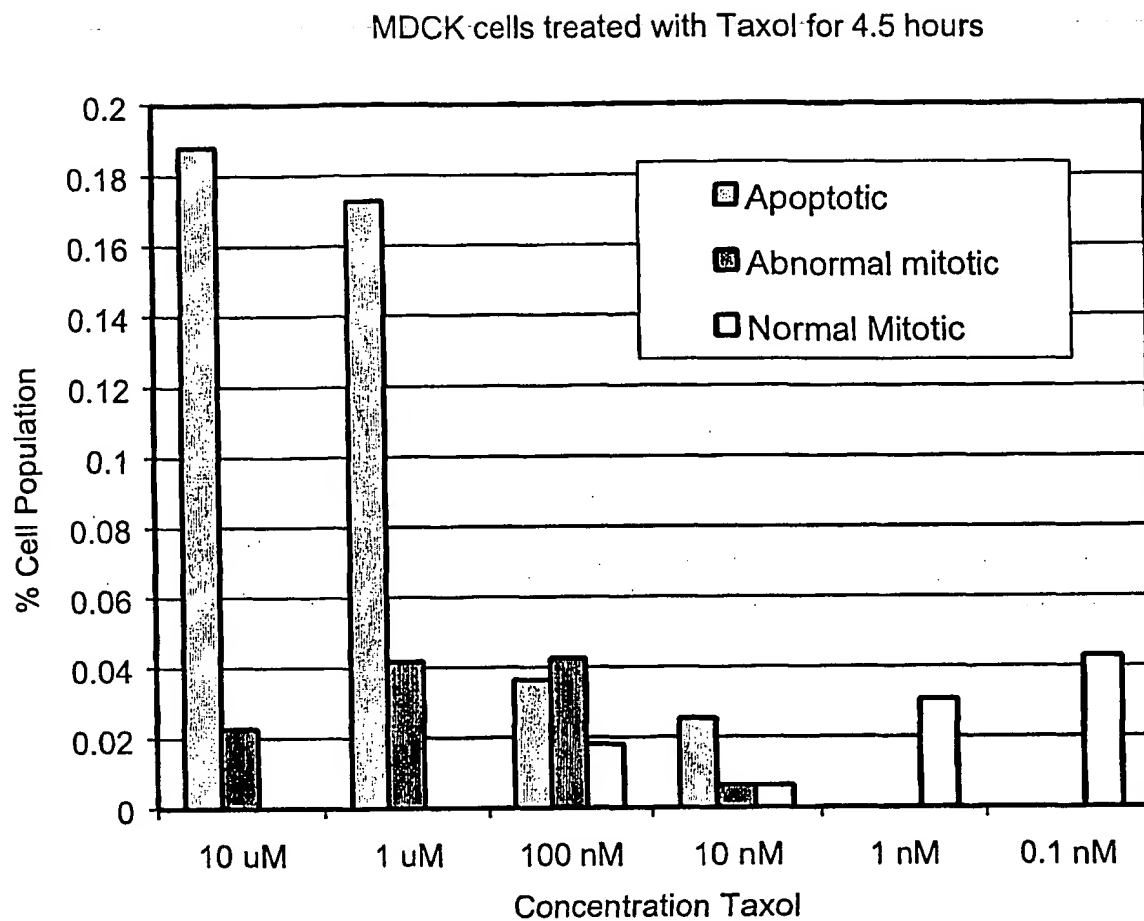


FIG. 10

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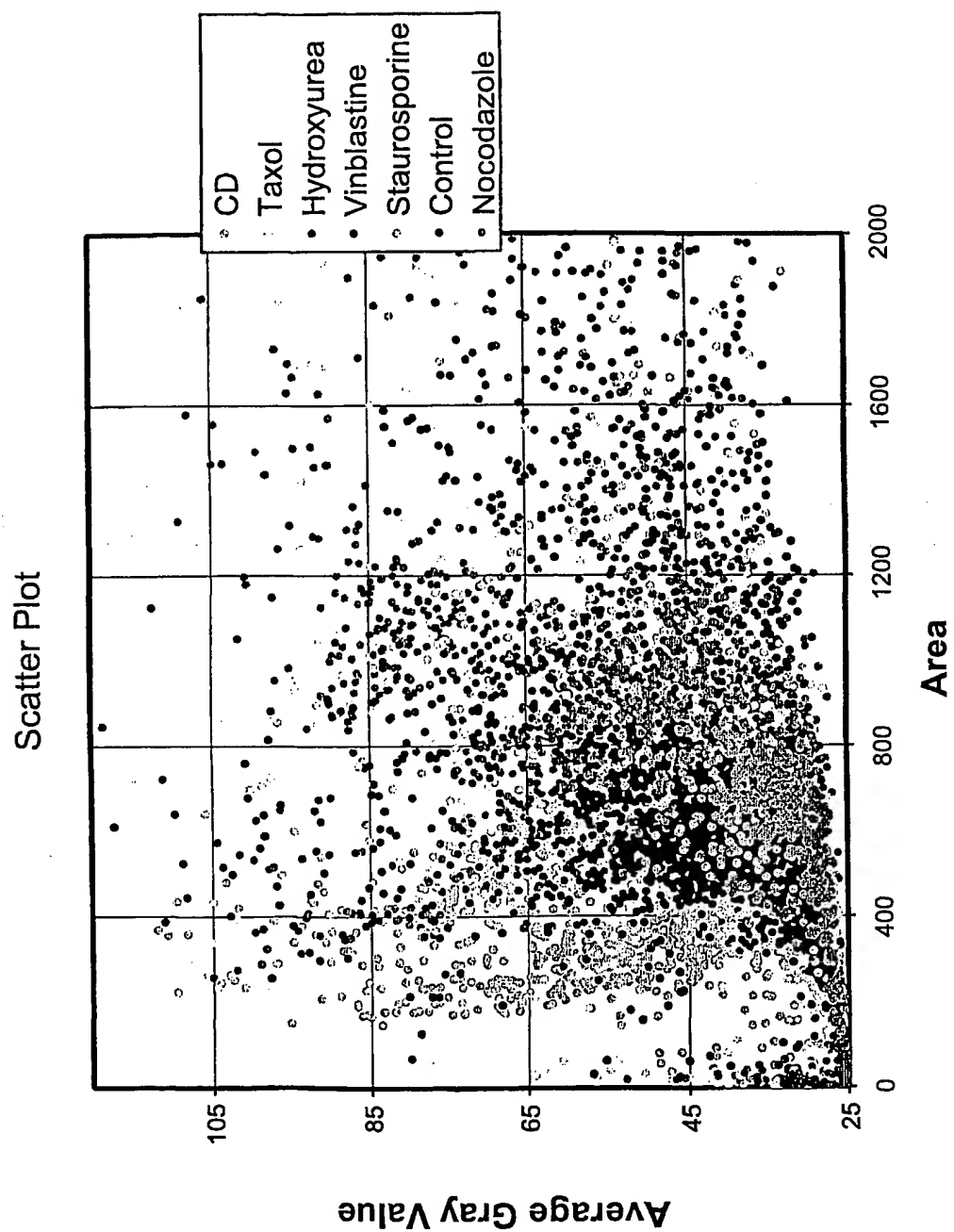


FIG. 11

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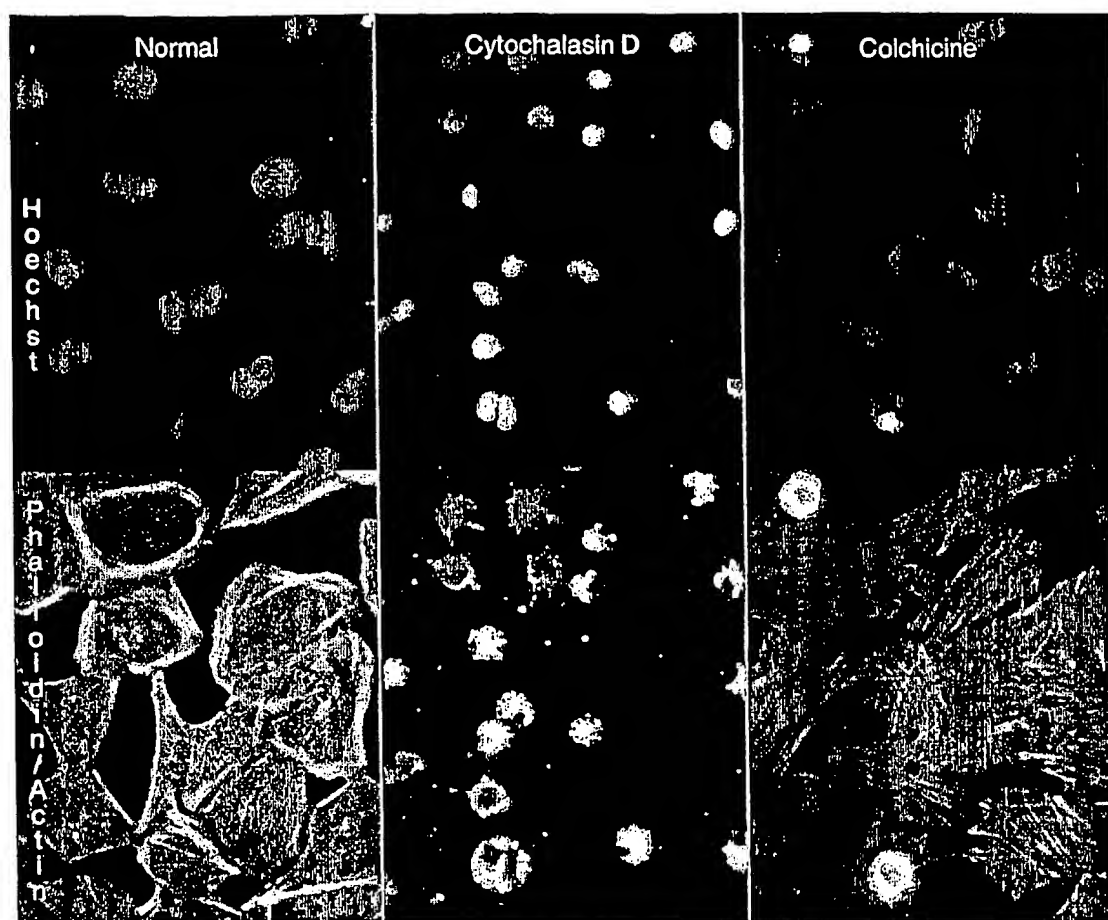


FIG. 12

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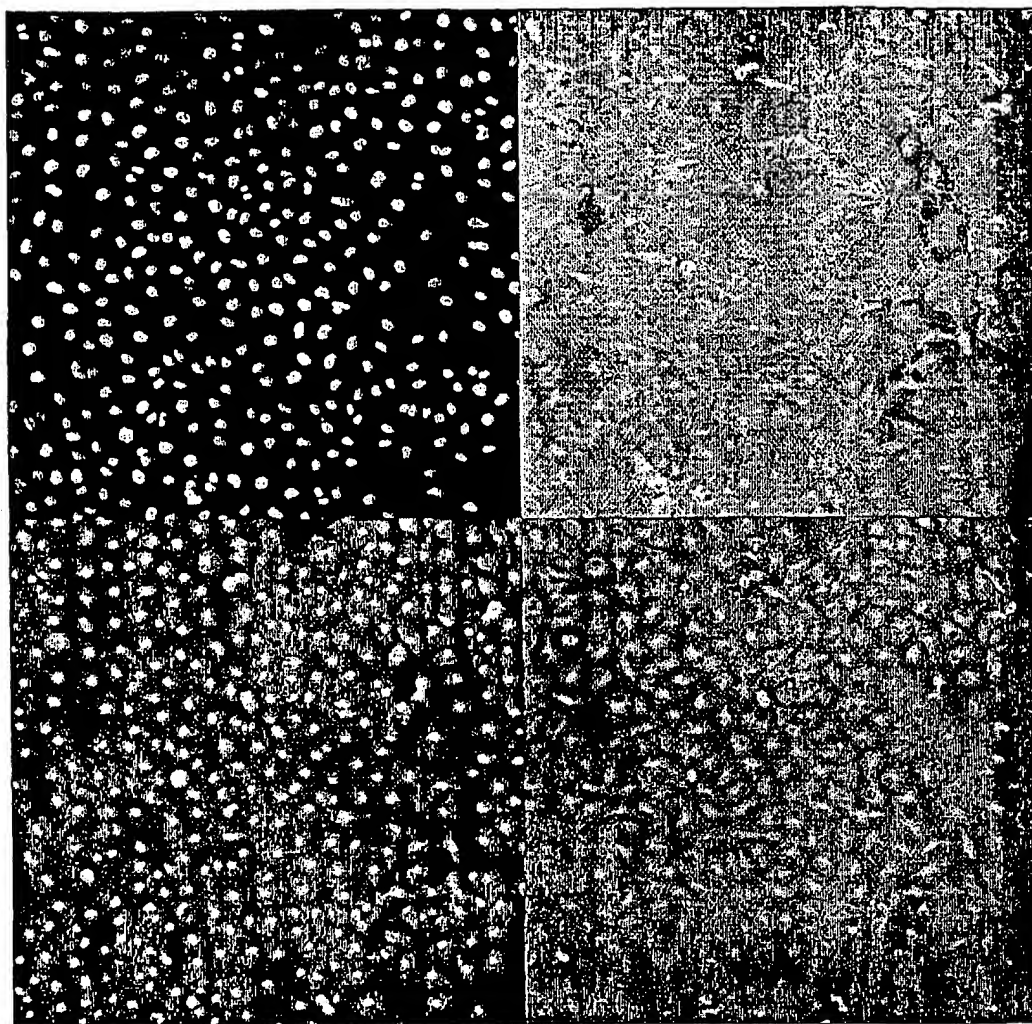


FIG. 13

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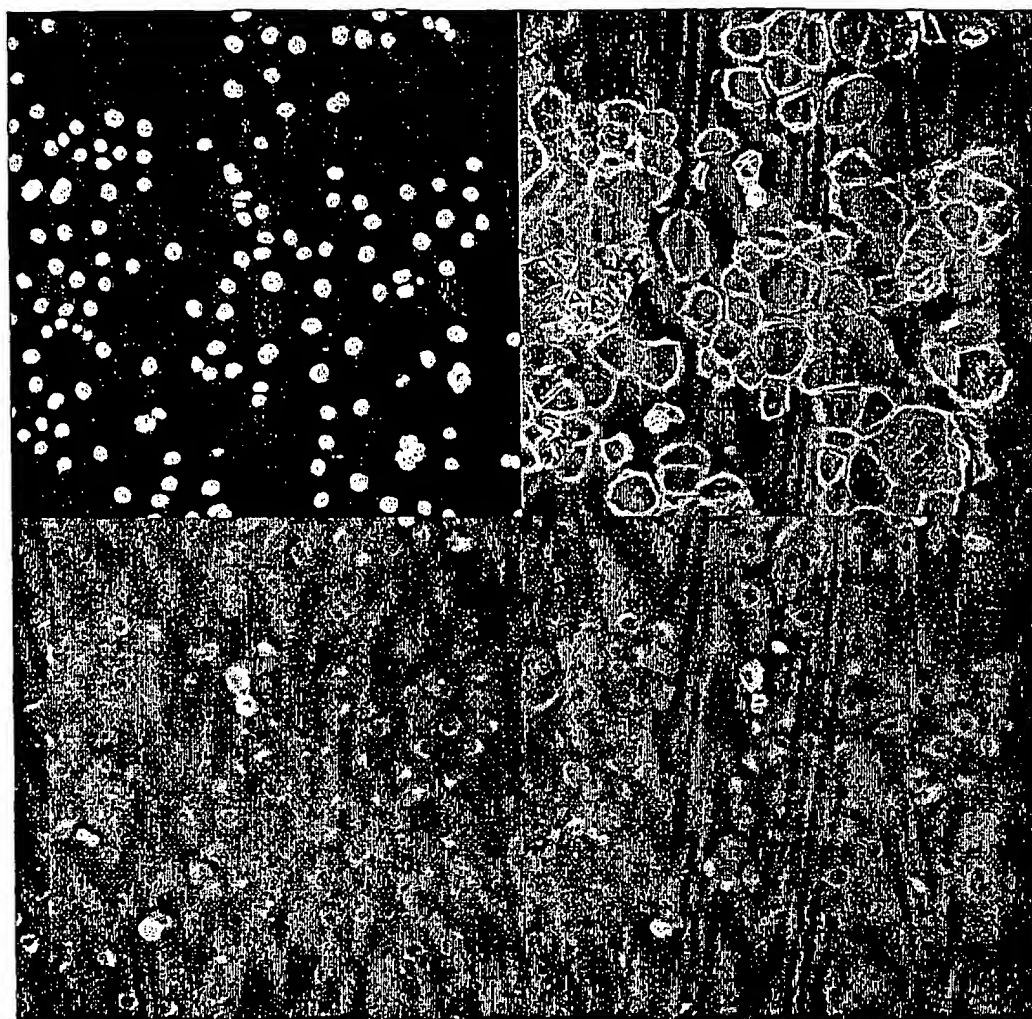


FIG. 14

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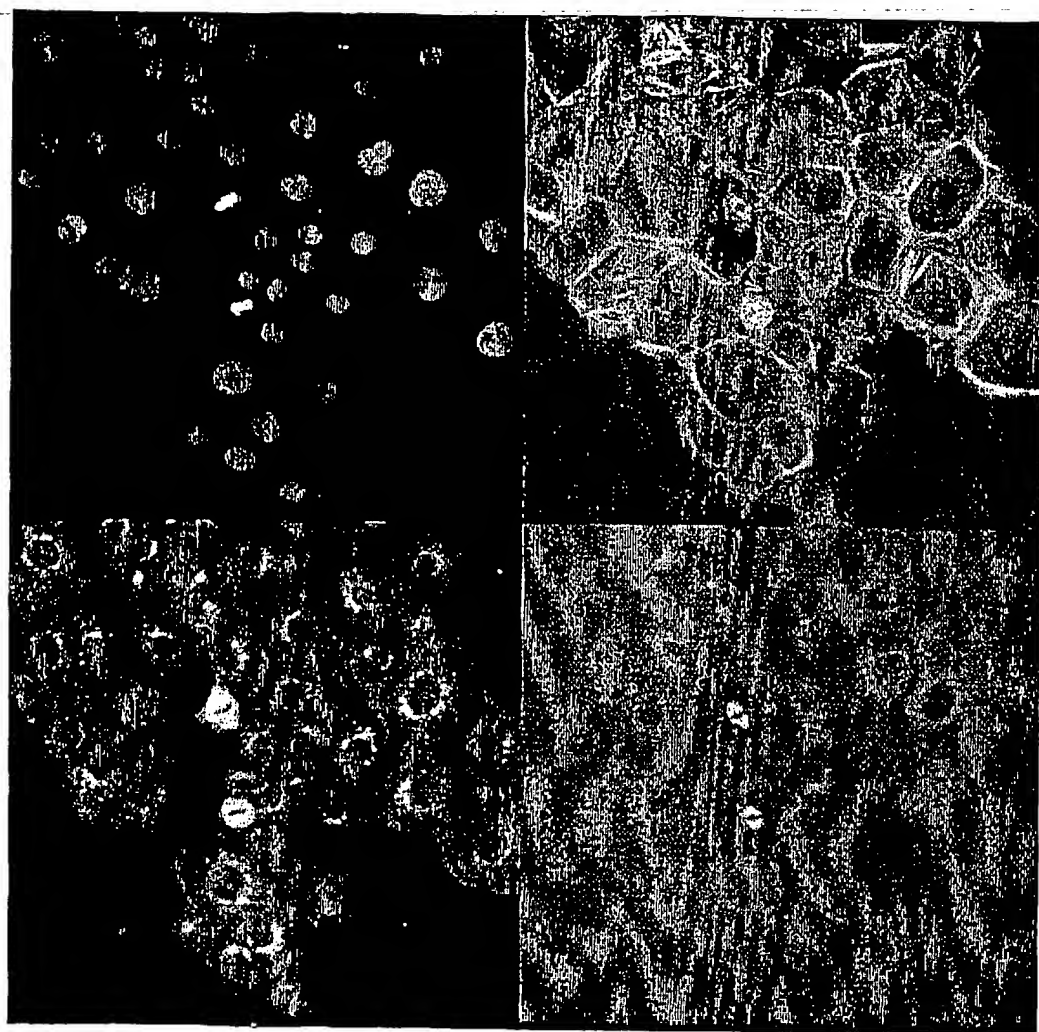


FIG. 15

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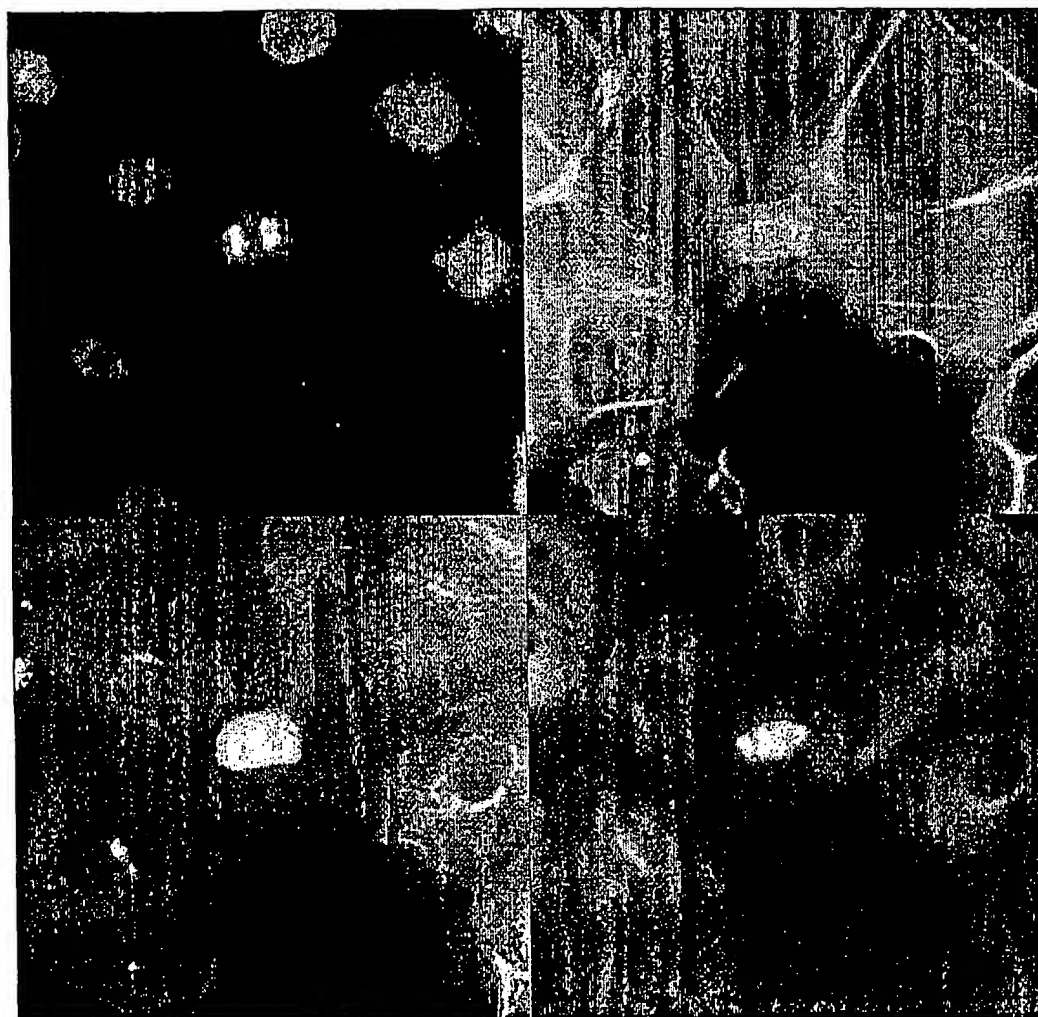


FIG. 16

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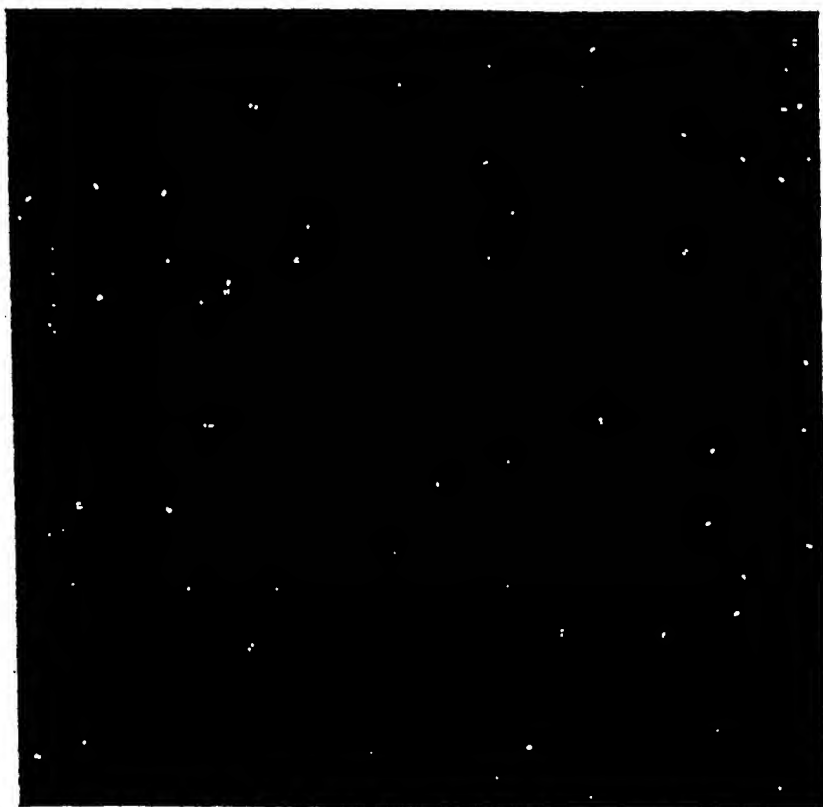


FIG. 17

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Conversion of morphometric parameters into nucleic acid code and clustering of the resulting sequences using Neighbor Joining method.

Compound:	Measurements																			
	Count	Area	Perimeter	Length	Breadth	Fiber length	Fiber breadth	Shape factor	Ell. form factor	Inner radius	Outer radius	Mean radius	Equiv. radius	Equiv. sphere vol.	Equiv. prolate vol.	Equiv. oblate vol.	Equiv. sphere surface area	Average gray value	Total gray value	Optical density
Control	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t
Taxol	a	t	t	t	t	t	t	t	a	t	t	t	t	t	t	t	t	t	t	t
CD	c	a	a	a	t	a	t	t	c	a	a	a	a	a	a	a	a	t	a	a
Nocodozol	c	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t
Staurosporine	g	g	c	a	a	t	a	a	t	g	a	a	a	t	g	g	g	a	a	t
Vinblastine	c	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	t	g	t	t
Hydroxyurea	g	t	t	t	t	t	t	g	t	t	t	t	t	t	t	t	t	t	c	t

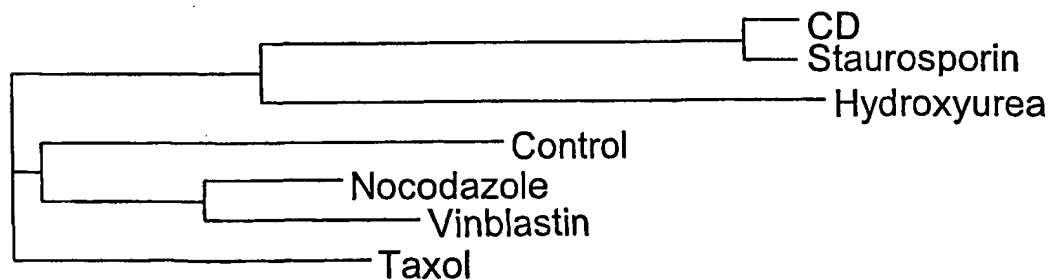


FIG. 18

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Conversion of morphometric parameters into amino acid codes and clustering of the resulting sequences using Neighbor Joining method.

	Count	Area	Perimeter	Length	Breadth	Fiber length	Fiber breadth	Shape factor	Ell. form factor	Inner radius	Outer radius	Mean radius	Equiv. radius	Equiv. sphere vol.	Equiv. prolate vol.	Equiv. oblate vol.	Equiv. sphere surface area	Average gray value	Total gray value	Optical density	Radial dispersion	Texture Difference Moment	EFA Harmonic 2, Semi-Major Axis	EFA Harmonic 2, Semi-Minor Axis	EFA Harmonic 2, Semi-Major A
Control	H	P	T	T	N	S	D	W	E	S	T	T	T	F	C	C	P	P	M	C	T	G	T	T	Y
Taxol	G	F	M	M	P	M	P	H	G	S	M	M	W	C	F	P	F	R	C	M	M	H	M	P	S
CD	F	G	G	G	M	G	M	K	A	G	G	G	G	G	G	G	G	H	G	G	G	M	G	V	H
Nocodazol	W	F	M	M	W	M	P	T	R	S	M	M	M	F	M	W	F	M	M	R	M	M	M	F	G
Staurosporine	N	V	A	G	G	M	G	G	Y	V	G	G	G	M	V	V	V	G	G	H	G	M	G	G	V
Vinblastine	F	W	W	M	W	W	C	W	D	S	M	W	W	M	M	M	W	M	V	E	M	M	M	F	P
Hydroxyurea	S	H	H	H	H	H	H	V	H	H	H	H	H	H	H	H	H	H	H	A	H	G	H	H	D

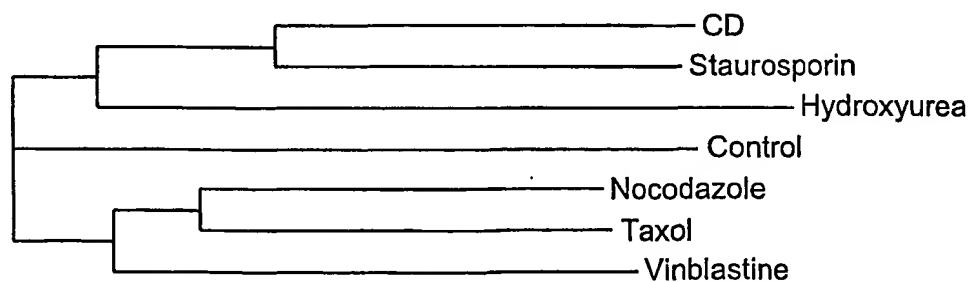


FIG. 19

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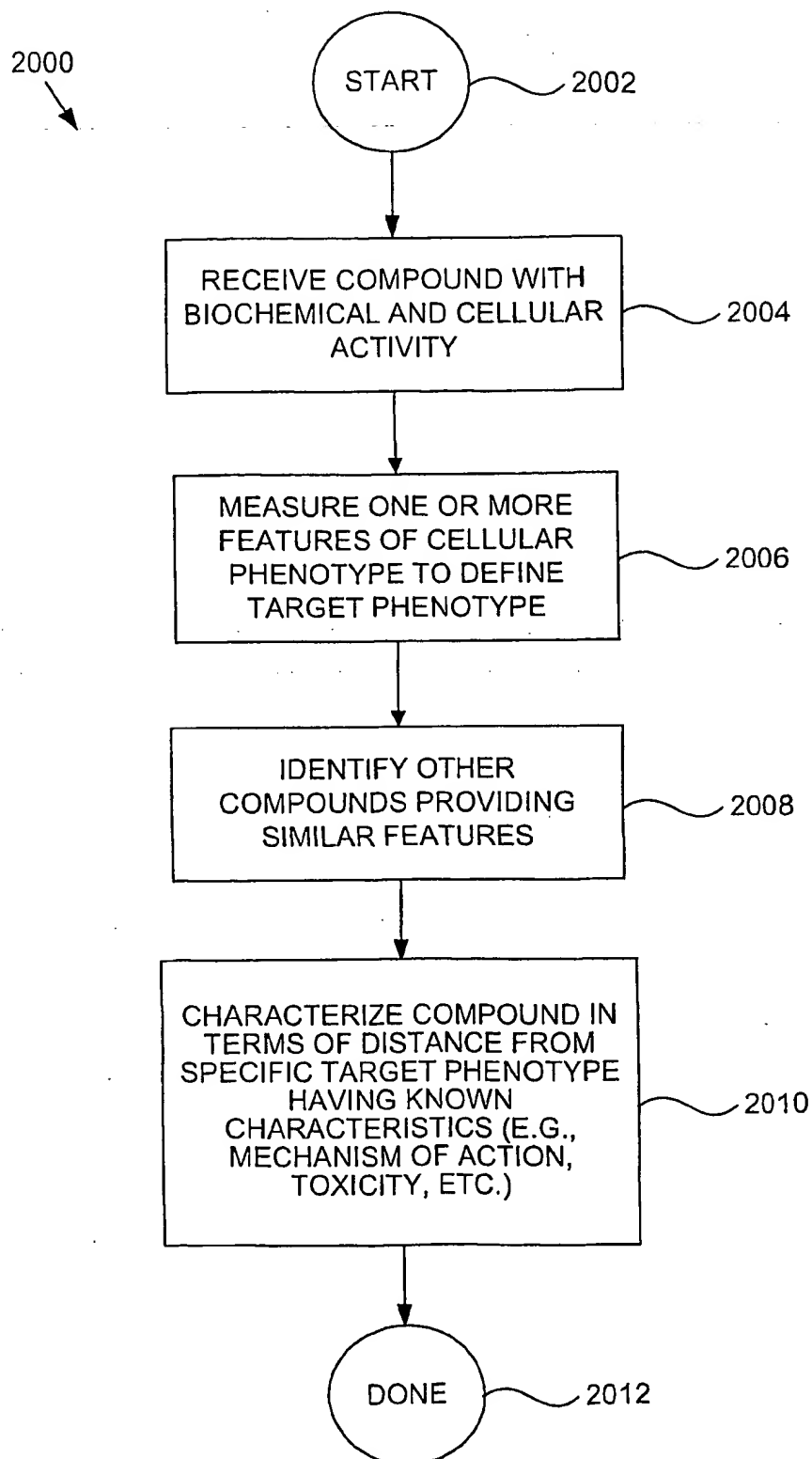


FIG. 20

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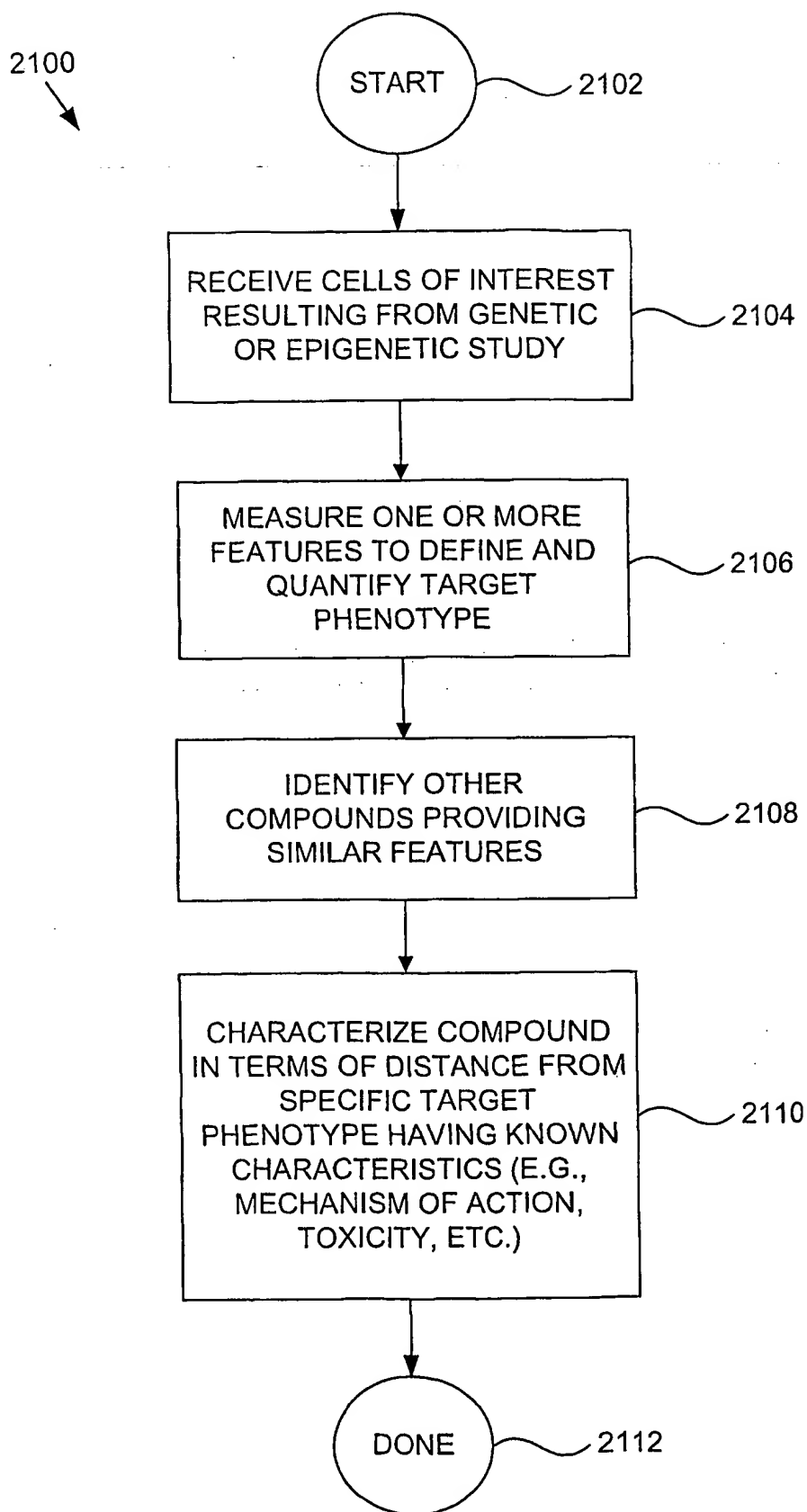
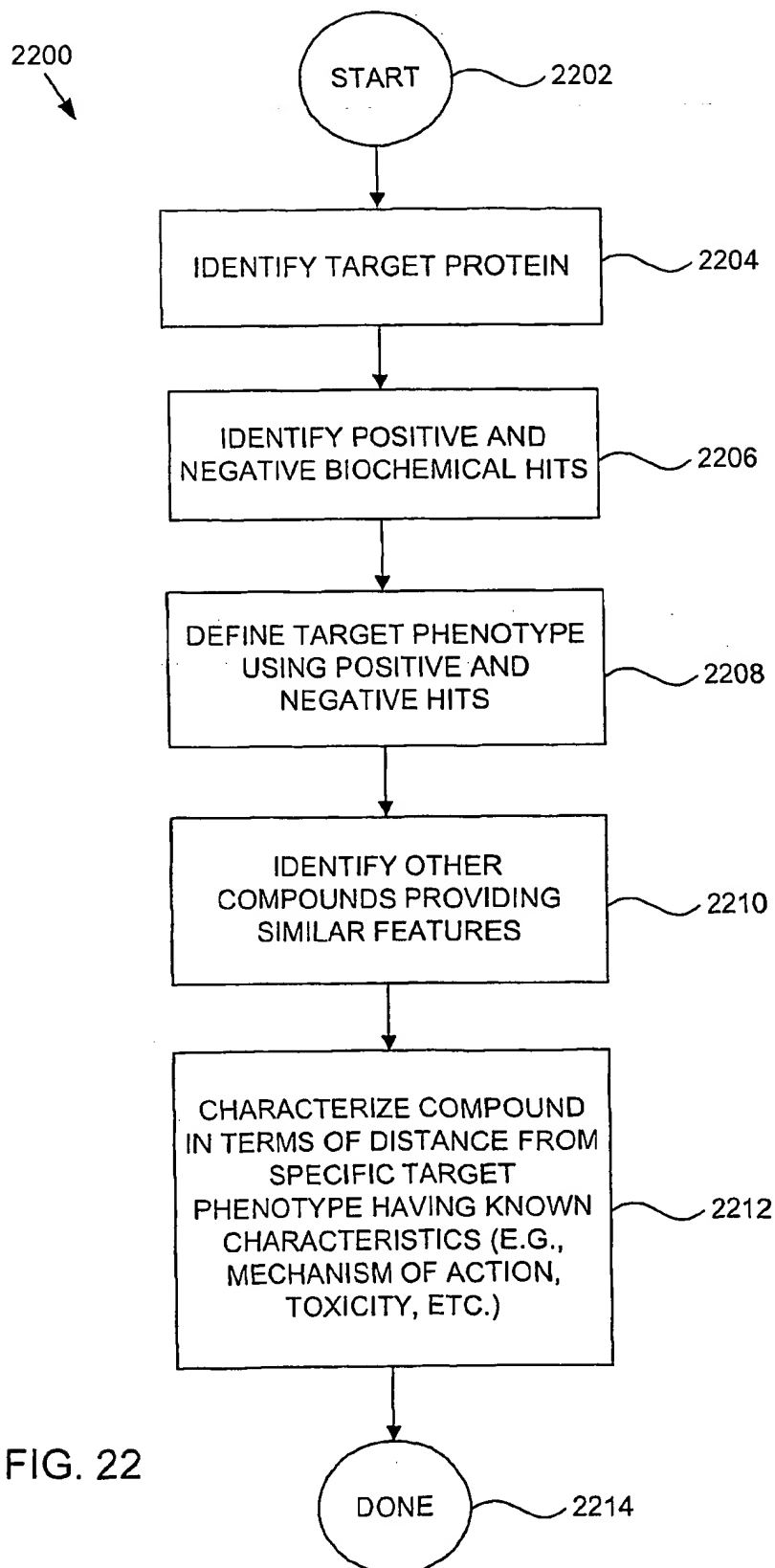


FIG. 21

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INTERNATIONAL SEARCH REPORT

Intel. .onal Application No

PCT/US 00/13154

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 G06F19/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 G06F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 98 38490 A (BIODX INC ;DUNLAY R TERRY (US); GOUGH ALBERT H (US); GIULIANO KENN) 3 September 1998 (1998-09-03) cited in the application	1-6, 24-27
Y	page 1; claims 1-43	7-23
X	WO 98 45704 A (TULLIN SOEREN ;KASPER ALMHOLT (DK); NOVONORDISK AS (DK); SCUDDER K) 15 October 1998 (1998-10-15) abstract; claims 1-3,22,73,80,81,86	1-6, 24-27
	-/--	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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- *O* document referring to an oral disclosure, use, exhibition or other means
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- *&* document member of the same patent family

Date of the actual completion of the international search

17 November 2000

Date of mailing of the international search report

24/11/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax (+31-70) 340-3000

Authorized officer

Filloy García, E

INTERNATIONAL SEARCH REPORT

Inter. Application No

PCT/US 00/13154

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	MONTIRONI R ET AL: "COMPUTED CELL CYCLE AND DNA HISTOGRAM ANALYSES IN IMAGE CYTOMETRY IN BREAST CANCER" JOURNAL OF CLINICAL PATHOLOGY, GB, LONDON, vol. 46, no. 9, 1 September 1993 (1993-09-01), pages 795-800, XP000644549 ISSN: 0021-9746 abstract ----	7-13
Y	WO 97 40055 A (DOW CHEMICAL CO ;UNIV TEXAS TECH (US)) 30 October 1997 (1997-10-30) page 18, line 26 - line 32 ----	14-23
P,X	WO 99 39184 A (HARTMANN THOMAS ;RIBOZYME PHARM INC (US)) 5 August 1999 (1999-08-05) the whole document ----	1-6, 24-27
P,X	WO 00 17643 A (CELLOMICS INC ;DUNLAY R TERRY (US); GOUGH ALBERT H (US); RUBIN RIC) 30 March 2000 (2000-03-30) the whole document ----	1-6, 24-27
E	WO 00 50872 A (CELLOMICS INC ;KAPUR RAVI (US); GIULIANO KENNETH A (US)) 31 August 2000 (2000-08-31) the whole document ----	1-6, 24-27
A	GIULIANO K A ET AL: "Fluorescent-protein biosensors: new tools for drug discovery" TRENDS IN BIOTECHNOLOGY, GB, ELSEVIER PUBLICATIONS, CAMBRIDGE, vol. 16, no. 3, 1 March 1998 (1998-03-01), pages 135-140, XP004108592 ISSN: 0167-7799 page 139, left-hand column, paragraph 4 -right-hand column, paragraph 3 -----	1-27